



THE BRITISH LIDITARY SCIENCYORED INTELLECTUAL PROPERTY ORGANIZATION International Bureau THE BRITISH LIDITARY



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 5: C12N 15/12, 15/54, 9/12 // C12Q 1/68, C12N 15/11

A1 (11) International Publication Number:

WO 93/15201

(43) International Publication Date:

5 August 1993 (05.08.93)

(21) International Application Number:

PCT/US93/00586

(22) International Filing Date:

22 January 1993 (22.01.93)

(30) Priority data:

826,935

22 January 1992 (22.01.92) US

(60) Parent Application or Grant (63) Related by Continuation

US Filed on

826,935 (CIP) 22 January 1992 (22.01.92)

(71) Applicant (for all designated States except US): NEW ENG-LAND DEACONESS HOSPITAL [US/US]; 185 Pilgrim Road, Boston, MA 02215 (US). (72) Inventors; and

(75) Inventors/Applicants (for US only): AVRAHAM, Hava [IL/US]; 50 Radmor Road, Brighton, MA 02135 (US). GROOPMAN, Jerome [US/US]; 79 Druce Street, Brookline, MA 02146 (US). COWLEY, Sally [GB/GB]; 28 Avonmore Road, London W14 8RS (GB). SCAD-DEN, David [US/US]; 62 Lexington Street, Weston, MA 02193 (US).

(74) Agents: GRANAHAN, Patricia et al.; Hamilton, Brook, Smith & Reynolds, Two Militia Drive, Lexington, MA + 02173 (US).

(81) Designated States: AU. CA. JP. US. European patent (AT. BE, CH, DE, DK, ES, FR, GB, GR. IE, IT, LU, MC. NL, PT, SE).

Published

With international search report. With amended claims.

Date of publication of the amended claims:

30 September 1993 (30.09.93)

(54) Title: NOVEL PROTEIN TYROSINE KINASES

(57) Abstract

The identification and isolation of novel protein tyrosine kinase genes present on human megakaryocytic and lymphocytic cells, the proteins encoded by these genes, antibodies specific for the encoded proteins, RNA nucleic acid sequences which hybridize to the genes and methods of use therefor.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AT	Austria	FR	France	MR	Mauritania
AU	Australia	GA	Gabon	MW	Malawi
88	Barhados	GB	United Kingdom	NL	Netherlands
BE	Belgium	GN	Guinea	NO	Norway
BF	Burkina Faso	GR	Greece	NZ	New Zealand
BC	Bulgaria	HU	Hungary	PL	Poland
BJ	Benin	ΙE	Ireland	PT	Portugal
BR	Brazil	IT	Italy	RO	Romania
CA	Canada	JP	Japan	RU	Russian Federation
CF	Central African Republic	KP	Democratic People's Republic	SU	Sudan
CC	Congo		of Korea	SE	Sweden
СН	Swiperland	KR	Republic of Korea	SK	Slovak Republic
CI	Côte d'Ivoire	ĸz	Kiwakhstan	SN	Senegal
CM	Canteroon	1.1	Lieghtenstein	SU	Soviet Union
CS	Częchosłowakia	LK	Sri Lanka	TD	Chud
CZ	Czech Republic	1.13	Luxembourg	TG	Togo
DΕ	Germany	M(C	Monaco	UA	Ukraine
DK	Denmark	MC	Madagascar	us	United States of America
ES	Spain	MI	Mali	VN	Viet Nam
FI	Finland	MN	Mongolia		

NOVEL PROTEIN TYROSINE KINASES

Description

Background of the Invention

Transduction of signals that regulate cell growth and differentiation is regulated in part by phosphorylation of various cellular proteins. Protein tyrosine kinases are enzymes that catalyze this process. Moreover, many act as growth factor receptors.

Summary of the Invention

The present invention relates to novel protein tyrosine kinase genes present in human megakaryocytic and lymphocytic cells, the proteins encoded by these genes, antibodies specific for the encoded proteins, RNA nucleic acid sequences which hybridize to the genes and methods of use therefor.

The genes isolated as described herein are referred to, collectively, as protein tyrosine kinase (pTK) genes. The nucleic acid sequences of these genes, isolated as discussed herein, show significant homology with previously identified protein tyrosine kinases containing extracellular domains which function as growth factor receptors. The pTK genes have been shown to be present in both megakaryocytic and lymphocytic cells.

The pTK genes of the present invention show

25 significant sequence homology with members of the c-kit subgroup of growth factor receptors with protein tyrosine kinase activity. The c-kit subgroup of receptor tyrosine kinases catalyze the phosphorylation of exogenous substrates, as well as tyrosine residues within their own polypeptide chains. (Ullrich, A. and Schlessinger, J., Cell, 61:203 (1990)). Members of the c-kit subgroup include FLT/FLK (Fetal Liver Kinase), FGF (Fibroblast)

Growth Factor Receptor) and NGF (Nerve Growth Factor Receptor).

In particular, fourteen pTK genes have been identified. Two pTK genes, referred to as SAL-S1 and SAL-5 D4 (also referred to as megakaryocyte derived FGF-like receptor tyrosine kinase) were identified in megakaryocytic cells. Five pTK genes, referred to as LpTKs, were identified in lymphocytic cells and have been shown to be present in megakaryocytes as well. One pTK gene, referred to a HpTKs, was identified in human hepatoma cells. Six pTK genes, referred to as bpTK genes, found in human brain tissue.

SAL-S1 is related to the FLT/FLK family of pTKs.

SAL-D4 is related to the FGF receptor family of pTKs, and

one LpTK (LpTK 3) is related to the NGF receptor family of pTKs.

The pTK genes, which are the subject of the present invention, were identified using two sets of degenerative oligonucleotide primers: a first set which amplifies all pTK DNA segments (SEQ ID NOS:1-2), and a second set which amplifies highly conserved sequences present in the catalytic domain of the c-kit subgroup of pTKs (SEQ ID NOS:3-4). The pTK genes identified in this manner are described below.

25 SAL-S1 is expressed in several megakaryocytic cell lines, but not in erythroid cell lines. The nucleotide sequence of SAL-S1 was obtained, revealing a sequence containing 158 base pairs. (SEQ ID NO:5). This isolated DNA fragment encoded an amino acid sequence (SEQ ID NO:6) which exhibited significant sequence homology with known protein tyrosine kinases of the FLT/FLK family. The full

length gene sequence (SEQ ID NO: 17) contains 6827 b.p. and the deduced amino acid sequence (SEQ ID NO: 18) contains 349 residues.

SAL-D4, also expressed in megakaryocytic cells, is a DNA fragment containing the nucleotide sequence of 141 base pairs. (SEQ ID NO:7). This isolated DNA fragment encoded an amino acid sequence (SEQ ID NO:8) which exhibited significant sequence homology with known protein tyrosine kinases of the FGF receptor family.

The LpTKs, including LpTK 2, LpTK 3, LpTK 4, and LpTK 13 and LpTK 25, are expressed in lymphocytic cells, as well as megakaryocytic cells. The nucleotide sequence (151 base pairs) of the LpTK 3 gene was obtained (SEQ ID NO:11), and exhibited significant homology with known

protein tyrosine kinases of the NGF receptor family. The nucleotide sequences of the LpTK 2, LpTK 4, and LpTK 13 genes contained 149 base pairs (SEQ ID NO:9), 137 base pairs (SEQ ID NO:13), and 211 base pairs (SEQ ID NO:15) respectively. LpTK 25 has a nucleotide sequence of 3120

b.p. (SEQ ID NO: 22). A full length gene sequence has been obtained for LpTK 2 (SEQ ID NO: 19) which contains 7606 b.p. Additional sequencing of LpTK 4 revealed a sequence of 404 b.p. (SEQ ID NO: 21).

The HpTK 5 gene, expressed in human hepatoma cells,

25 has a nucleotide sequence of 3120 b.p. (SEQ ID NO: 22).

Nucleotide sequences of the bpTK's, including bpTK 1, bpTK

2, bpTK 3, bpTK 4, bpTK 5 and bpTK 7 are expressed in

human brain tissue encode proteins having the amino acid

sequences of SEQ ID NOS: 25-30 respectively.

Thus the present invention includes DNA isolated from a human megakaryocytic cell line, which hybridizes to a

DNA fragment which hybridizes to DNA encoding an amino acid sequence which is highly conserved in the catalytic domain of protein tyrosine kinases of the c-kit subgroup.

The present invention also includes the proteins 5 encoded by the pTK genes identified as described herein, which exhibit significant sequence homology with members of the c-kit subgroup of pTks (i.e. FLT/FLK (SAL-S1), FGF receptor (SAL-D4) or NGF receptor (LpTKS)) as well as the proteins encoded by HpTK 5 and the bpTKs. The present 10 invention also includes SAL-S1, SAL-D4, and LpTK, HpTK and bpTK homologues or equivalents (i.e., proteins which have amino acid sequences substantially similar, but not identical, to that of SAL-S1, SAL-D4, the LpTKs HpTK and the bpTKs, which exhibit tyrosine kinase activity.) 15 invention further includes peptides (SAL-S1, SAL-D4, LpTK, HpTK and bpTK fragments) which retain tyrosine kinase activity, yet are less than the entire SAL-S1, SAL-D4, LpTK, HpTK and bpTK sequences), monoclonal and polyclonal antibodies specific for SAL-S1, SAL-D4, the LpTKs, HpTK 20 and the bpTKs, and uses for the SAL-S1, SAL-D4, the LpTK, HpTK and the bpTK nucleic acid sequences and SAL-S1, SAL-D4, LpTK, HpTK and bpTK equivalents.

The present invention further includes nucleic acid sequences which hybridize with DNA or RNA encoding the proteins described herein, which exhibit significant sequence homology with the FLT/FLK, FGF receptor or NGF receptor family of protein tyrosine kinases contained within the c-kit subgroup. Such nucleic acid sequences are useful as probes to identify pTK genes in other vertebrates, particularly mammals, and in other cell types. They can also be used as anti-sense

oligonucleotides to inhibit protein tyrosine kinase activity, both in vitro and in vivo.

The SAL-S1, SAL-D4, LpTK, HpTK and bpTK, tyrosine kinases of the present invention can be used as target proteins in conjunction with the development of drugs and therapeutics to modulate cell growth, differentiation and other metabolic functions. The SAL-S1, SAL-D4, LpTK, HpTK or bpTK proteins can be used as agonists or antagonists to other tyrosine kinases. The SAL-S1, SAL-D4, LpTK, HpTK or bpTK tyrosine kinases can also be instrumental in the modulation of megakaryocyte and/or platelet adhesion interactions.

In addition, the SAL-S1, SAL-D4, LpTK, HpTK and bpTK tyrosine kinases can be used in screening assays to detect cellular growth and/or differentiation factors. Using standard laboratory techniques, the ligands of the pTKs of the present invention can be identified. Once identified, assays can be designed to detect these ligands present endogenously, within cells, as well as exogenously, in extra cellular fluids. Assays can also be designed as diagnostic aids to detect these ligands in body fluids such as blood and urine.

Brief Description of the Drawings

Figure 1 depicts the nucleotide sequence of SAL-S1 (SEQ ID NO: 5) and the deduced amino acid sequence (SEQ ID NO:6).

Figure 2 depicts the nucleotide sequence of SAL-D4 (SEQ ID NO:7) and its deduced amino acid sequence (SEQ ID NO:8).

Figure 3A depicts the nucleotide sequence (SEQ ID NO:9) and its deduced amino acid sequence (SEQ ID NO:10) for LpTK 2.

Figure 3B depicts the nucleotide sequence (SEQ ID NO:11) and its deduced amino acid sequence (SEQ ID NO:12) for LpTK 3.

Figure 3C depicts the nucleotide sequence (SEQ ID NO:13) and its deduced amino acid sequence (SEQ ID NO:14) for LpTK 4.

Figure 3D depicts the nucleotide sequence (SEQ ID NO:15) and its deduced amino acid sequence (SEQ ID NO:16) for the LpTK 13.

Figure 4A-4J depicts the full-length nucleotide sequence (SEQ ID NO: 17) and its deduced amino acid sequence (SEQ ID NO: 18) for SAL-S1.

Figure 5A-5J depicts the full length nucleotide sequence (SEQ ID NO: 19) and the deduced amino acid sequence (SEQ ID NO: 20) for LpTK2.

Figure 6 depicts the partial nucleotide sequence (SEQ 20 ID NO: 21) for LpTK4.

Figure 7A-7D depicts the full length nucleotide sequence (SEQ ID NO: 22) for LpTK25.

Figure 8A-8F depicts the full length nucleotide sequence (SEQ ID NO: 23) and the deduced amino acid sequence (SEQ ID NO: 24) for HpTK5.

Figure 9 depicts the amino acid sequence (SEQ ID NO:

25) of bpTK1.

Figure 10 depicts the amino acid sequence (SEQ ID NO:

- 26) of bpTK2.
- Figure 11 depicts the amino acid sequence (SEQ ID NO: 27) of bpTK3.

Figure 12 depicts the amino acid sequence (SEQ ID NO: 28) of bpTK4.

Figure 13 depicts the amino acid sequence (SEQ ID NO: 29) of bpTK5.

Figure 14 depicts the amino acid sequence (SEQ ID NO: 30) of bpTK7.

Detailed Description of the Invention

Novel protein tyrosine kinase genes have been identified, their nucleic acid sequences determined, and the amino acid sequences of the encoded proteins deduced. The genes isolated as described herein are referred to, collectively, as protein tyrosine kinase (pTK) genes. The nucleic acid sequences of these genes, isolated as discussed herein, show significant homology to with previously identified protein tyrosine kinases containing extracellular domains which function as growth factor receptors. These genes have been shown to be present in both megakaryocytic and lymphocytic cells.

To facilitate the isolation and identification of
these novel pTKs, two sets of DNA probes were used, as
described in the Exemplification. The first set consisted
of two degenerative oligonucleotide sequences, pTK 1 (SEQ
ID NO:1) and pTK 2 (SEQ ID NO:2) (Matthews, W. Cell 65:
1143 (1991; Wilks, A. F. Proc. Natl. Acad. Sci. USA
25 86:1603 (1989)). These sequences were used as primers in
a polymerase chain reaction to amplify tyrosine kinase DNA
segments. (Mullis, K. et al., Cold Spring Harbor Symp.
Advan. Biol. 51:263 (1986).

The second set consisted of two oligonucleotide 30 sequences, pTK 3 (SEQ ID NO:3) and pTKKW (SEQ ID NO:4)

designed to amplify the nucleic acid sequence which encodes the highly conserved regions of the catalytic domains of the c-kit family of protein tyrosine kinases. These sequences were used as primers in the polymerase chain reaction in a second round of DNA amplification. Using this two-step amplification procedure, DNA fragments which hybridized to these pTK primers were identified, isolated and subsequently sequenced.

In particular, fourteen pTK genes exhibiting

10 significant homology with the c-kit subgroup of protein tyrosine kinases have been identified. Two pTK genes, referred to as SAL-S1 and SAL-D4 (also referred to as megakaryocyte derived FGF-like receptor) were identified in several megakaryocytic cell lines, including CMK 11-5,

15 DAMI, UT-7 and UT-7 grown in erythropoietin, but not in the erythroid cell lines HEL, PMA stimulated HEL cells, or K562. Five pTK genes, referred to as LpTKs, were identified in lymphocytic, as well as in megakaryocytic cells. One pTK gene, referred to as HpTK5 was identified in human hepatoma cells and six genes, referred to as bpTKs, were identified in human brain tissue.

SAL-S1 (SEQ ID NO:6 and 18) encoded by the nucleic acid sequence SEQ ID NOS:5 and 17, exhibits significant homology with the FLT/FLK family of pTKs. SAL-D4 (SEQ ID NO:8) encoded by SEQ ID NO:7, is related to the FGF receptor family of pTKs, and one LpTK (LpTK 3 (SEQ ID NO:12) encoded by the SEQ ID NO:11, is related to the NGF receptor family of pTKs. The remaining LpTKs, LpTK2 (SEQ ID NO:10) encoded by SEQ ID NO:9; LpTK4 (SEQ ID NO:14) encoded by SEQ ID NO:13; LpTK13 (SEQ ID NO:16) encoded by SEQ ID NO:15 LpTK25 encoded by SEQ ID NO: 22, also exhibit

sequence homology with known protein tyrosine kinases (Data not shown).

HpTK5 (SEQ ID NO: 24) encoded by SEQ ID NO: 23 and the bpTKs 1, 2, 3, 4, 5 and 7 (SEQ ID NOS: 25-30 respectively, also exhibit sequence homology with known protein tyrosine kinases.

Thus, as described above, DNA which hybridize with DNA encoding amino acid sequences present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein kinases have been isolated and sequenced. These isolated DNA sequences, collectively referred to as pTKs genes, (and their deduced amino acid sequences) have been shown to exhibit significant sequence homology with

known members of receptor tyrosine kinase families.

Once isolated, these DNA fragments can be amplified using known standard techniques such as PCR. These amplified fragments can then be cloned into appropriate cloning vectors and their DNA sequences determined.

These DNA sequences can be excised from the cloning vectors, labeled with a radiolabeled nucleotide such as ³²P and used to screen appropriate cDNA libraries to obtain the full-length cDNA clone.

The pTk genes as described above have been isolated from the source in which they occur naturally, i.e.

25 megakaryocyte and lymphocytic cells. The present invention is intended to include pTk genes produced using genetic engineering techniques, such as recombinant technology, as well as pTk genes that are synthesized chemically.

The deduced amino acid sequences of the pTK genes

on include amino acid sequences which encode peptides

exhibiting significant homology with the catalytic domain

of protein tyrosine kinases of the c-kit subgroup of tyrosine kinases. These proteins, encoded by the pTk genes, can include sequences in which functionally equivalent amino acid residues are substituted for residues within the sequence, resulting in a silent change, that is a change not detected phenotypically. For example, one or more amino acid residues within the sequence can be substituted by another amino acid of a similar polarity which acts as a functional equivalent, resulting in a silent substitution.

In addition, the protein structure can be modified by deletions, additions, inversion, insertions or substitutions of one or more amino acid residues in the sequence which do not substantially detract from the desired functional tyrosine kinases properties of the peptide.

Modified pTKs of the present invention, with receptor tyrosine kinase activity can be made using recombinant DNA techniques, such as excising it from a vector containing a cDNA encoding such a protein, or by synthesizing DNA encoding the desired protein mechanically and/or chemically using known techniques.

An alternate approach to producing the pTKs of the present invention is to use peptide synthesis to make a peptide or polypeptide having the amino acid sequence of such a protein. The peptides or modified equivalents thereof, can be synthesized directly by standard solid or liquid phase chemistries for peptide synthesis.

Preferably, the pTKs of the present invention will be 30 produced by inserting DNA encoding the proteins into an appropriate vector/host system where it will be expressed.

The DNA sequences can be obtained from sources in which they occur naturally, can be chemically synthesized or can be produced using standard recombinant technology.

This invention also pertains to an expression vector comprising a pTK gene of the present invention, encoding for a protein which exhibits receptor tyrosine kinase activity.

The pTK genes of the present invention can be used for a number of diagnostic and therapeutic purposes. For example, the nucleic acid sequences of the pTK genes can be used as probes to identify other protein tyrosine kinases present in other cell types, including eukaryotic and prokaryotic cell types.

The nucleic acid sequences can be used to design
drugs that directly inhibit the kinase activity of protein
tyrosine kinases, or to design peptides that bind to the
catalytic domain of tyrosine kinases, thus inhibiting
their activity. These sequences can also be used to
design anti-sense nucleotides that can also inhibit, or
destroy, tyrosine kinase activity. Such inhibition of
tyrosine kinase activity would be desirable in
pathological states where decreased cellular proliferation
would be beneficial, such as leukemias or other
malignancies.

The nucleic acid sequences can also be used to design drugs, peptides or anti-sense nucleotides as above, but with enhancing, rather than inhibitory effects, on tyrosine kinases. Such enhanced tyrosine kinase activity would result in increasing the phosphorylation of substrates (exogenous, as well as endogenous tyrosine residues). Enhanced effects would be desirable in states

where increased cellular proliferation would be beneficial, such as anemias, bleeding disorders and during surgical procedures.

The pTK genes of the present invention can also be used to obtain soluble fragments of receptor tyrosine kinases, capable of binding their respective ligands (i.e. fibroblast growth factor).

pTK genes encoding soluble receptor tyrosine kinase fragments can be produced using recombinant DNA techniques or synthetically. In either case, the DNA obtained encodes a soluble pTK fragment which lacks a substantial portion of the hydrophobic transmembrane region to permit solubilization of the fragment.

These soluble pTK protein fragments can be introduced exogenously to act as competitors with the endogenous, membrane bound pTK for their respective ligands, thus inhibiting tyrosine kinase activity. Alternately, a modified soluble pTK protein fragment can be introduced which binds the ligand but does not activate kinase activity.

These soluble pTK protein fragments can also be used in binding assays to detect ligands such as growth and differentiation factors. Once these ligands are identified, they may be altered or modified to inhibit or enhance kinase activity. For example, the ligands may be modified or attached to substances that are toxic to the cell, such a ricin, thus destroying the target cell. The substance may be a super-activating substance which, after binding to the pTK, may substantially increase the kinase activity, or activate other growth factors.

pTk genes of the present invention would also be useful to develop diagnostic tools for in vitro screening assays for ligands such as growth factors or differentiation factors that inhibit or enhance kinase activity. The proteins encoded by the pTK genes can also be used in such assays, or as immunogens to produce monoclonal or polyclonal antibodies to be used in such assays.

Such antibodies can also be used in methods of treating conditions in which an individual would benefit therapeutically if protein tyrosine kinase activity could be modified, such as increasing platelet production in bleeding disorders.

The present invention will now be illustrated by the following Exemplification, which is not intended to be limiting in any way.

Exemplification: The Identification and Isolation of the pTK Genes

To facilitate the isolation and identification of these novel pTK genes, two sets of DNA probes were used. (See Table).

The first set consisted of two degenerative oligonucleotide sequences, pTK 1 (SEQ ID NO:1) and pTK 2(SEQ ID NO:2). These sequences were used as polymerase chain reaction (PCR) primers, using standard PCR techniques, to amplify tyrosine kinase DNA segments.

The second set consisted of two oligonucleotide sequences, pTK 3 (SEQ ID NO:3) and pTKKW (SEQ ID NO:4) selected from the highly conserved regions of the catalytic domains of the c-kit subgroup of protein

-14-

tyrosine kinases. These sequences were also used as polymerase chain reaction primers in a second round of DNA amplification. Using this two-step amplification procedure, DNA fragments which hybridized to these pTK primers were identified, isolated and subsequently sequenced using known laboratory techniques.

TABLE

First Round of Amplification

PTK1

10 CGGATCCACAGNGACCT

PTK2

GGAATTCCAAAGGACCAGACGTC

Second Round of Amplification

PTK3 (kit family specific)

15 CGGATCCATCCACAGAGATGT

PTKKW (kit family specific)
GGAATTCCTTCAGGAGCCATCCACTT

Equivalents

Those skilled in the art will recognize, or be able to ascertain using no more than routine experimentation, many equivalents to the specific embodiments of the invention described herein. Such equivalents are intended to be encompassed by the following claims.

-15-

CLAIMS

The invention claimed is:

5

- 1. Isolated DNA of human megakaryocytic origin which hybridizes to a DNA fragment which hybridizes to DNA encoding an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein tyrosine kinases.
- 2. Isolated DNA of Claim 1 having a nucleotide sequence selected from the group of nucleotide sequences consisting of:

```
a) SAL-S1 (SEQ ID NOS:5 and 7);
```

- b) SAL-D4 (SEQ ID NO:7);
- c) LpTK 2 (SEQ ID NOS:9 and 19);
- d) LpTK 3 (SEQ ID NO:11);
- e) LpTk 4 (SEQ ID NOS:13 and 21);
 - f) LpTK 13 (SEQ ID NO:15);
 - g) LpTK 25 (SEQ ID NO: 22 and
 - h(HpTK 5 (SEQ ID NO: 23).
- 3. Isolated DNA of Claim 1 which encodes an amino acid sequence selected from the group consisting of:

```
a) SAL-S1 (SEQ ID NOS:6 and 18);
```

- b) SAL-D4 (SEQ ID NO:8);
- c) LpTK 2 (SEQ ID NOS:10 and 20);
- d) LpTK 3 (SEQ ID NO:12);
- 25 e) LpTK 4 (SEQ ID NO:14);
 - f) LpTK 13 (SEQ ID NO:16);
 - g) HpTK 5 (SEQ ID NO:24);

```
h) bpTK 1 (SEQ ID NO:25);
i) bpTK 2 (SEQ ID NO:26);
j) bpTK 3 (SEQ ID NO:27);
k) bpTK 4 (SEQ ID NO:28);
l) bpTK 5 (SEQ ID NO:29); and
m) bpTK 7 (SEQ ID NO:30).
```

- 4. Isolated DNA of human megakaryocytic origin which comprises a DNA fragment whose sequence encodes an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein tyrosine kinases.
 - 5. Isolated DNA of Claim 4 which encodes an amino acid sequence selected from the group consisting of:

```
a) SAL-S1 (SEQ ID NO:6);
```

- b) SAL-D4 (SEQ ID NO:8);
 - c) LpTK 2 (SEQ ID NO:10);
 - d) LpTK 3 (SEQ ID NO:12);
 - e) LpTk 4 (SEQ ID NO:14); and
 - f) LpTK 13 (SEQ ID NO:16).
- 20 g) HpTK 5 (SEQ ID NO:24);
 - h) bpTK 1 (SEQ ID NO:25);
 - i) bpTK 2 (SEQ ID NO:26);
 - j) bpTK 3 (SEQ ID NO:27);
 - k) bpTK 4 (SEQ ID NO:28);
- 25 1) bpTK 5 (SEQ ID NO:29); and
 - m) bpTK 7 (SEQ ID NO:30).

- 6. A homogeneous protein of human megakaryocytic origin which includes an amino acid sequence exhibiting sequence homology with the catalytic domain of tyrosine kinases of the c-kit family.
- 5 7. A homogeneous protein of Claim 6 in which the amino acid sequence is selected from the group consisting of:
 - a) SAL-S1 (SEQ ID NO:6);
 - b) SAL-D4 (SEQ ID NO:8);
- 10 c) LpTK 2 (SEQ ID NO:10);
 - d) LpTK 3 (SEQ ID NO:12);
 - e) LpTk 4 (SEQ ID NO:14); and
 - f) LpTK 13 (SEQ ID NO:16).
 - g) HpTK 5 (SEQ ID NO:24);
- 15 h) bpTK 1 (SEQ ID NO:25);
 - i) bpTK 2 (SEQ ID NO:26);
 - j) bpTK 3 (SEQ ID NO:27);
 - k) bpTK 4 (SEQ ID NO:28);
 - 1) bpTK 5 (SEQ ID NO:29); and
- 20 m) bpTK 7 (SEQ ID NO:30).
 - 8. A protein of human megakaryocytic origin which exhibits significant sequence homology with the FLT/FLK family of protein tyrosine kinases.
- 9. A protein of Claim 8 encoded by the nucleotide sequence (SEQ ID NO:5).
 - 10. A protein of Claim 8 encoded by the amino acid sequence (SEQ ID NO:6).

- 19. A DNA expression vector containing a DNA sequence of human megakaryocytic origin which hybridizes to a DNA fragment which hybridizes to DNA encoding an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein kinases.
- 20. The DNA expression vector of Claim 17 containing a DNA sequence selected from the group consisting of:
 - a) SAL-S1 (SEQ ID NO:5);
- 10 b) SAL-D4 (SEQ ID NO:7);
 - c) LpTK 2 (SEQ ID NO:9);
 - d) LpTK 3 (SEQ ID NO:11);
 - e) LpTk 4 (SEQ ID NO:13); and
 - f) LpTK 13 (SEQ ID NO:15).
- 15 g) LpTK 25 (SEQ ID NO: 22 and
 - h (HpTK 5 (SEQ ID NO: 23).
 - 21. A cell transformed by the expression vector of Claim 17.

AMENDED CLAIMS

[received by the International Bureau on 31 August 1993 (31.08.93); original claims 1, 2, 4-7 and 14-21 amended; remaining claims unchanged (5 pages)]

- 1. Isolated DNA of human origin which hybridizes to a DNA fragment which hybridizes to DNA encoding an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein tyrosine kinases.
- Isolated DNA of Claim 1 having a nucleotide sequence selected from the group of nucleotide sequences
 consisting of:
 - a) SAL-S1 (SEQ ID NOS:5 and 17);
 - b) SAL-D4 (SEQ ID NO:7);
 - c) LpTK 2 (SEQ ID NOS:9 and 19);
 - d) LpTK 3 (SEQ ID NO:11);
- e) LpTk 4 (SEQ ID NOS:13 and 21);
 - f) LpTK 13 (SEQ ID NO:15);
 - g) LpTK 25 (SEQ ID NO: 22); and
 - h) HpTK 5 (SEQ ID NO: 23).
- 3. Isolated DNA of Claim 1 which encodes an amino acid sequence selected from the group consisting of:
 - a) SAL-S1 (SEQ ID NOS:6 and 18);
 - b) SAL-D4 (SEQ ID NO:8);
 - c) LpTK 2 (SEQ ID NOS:10 and 20);
 - d) LpTK 3 (SEQ ID NO:12);
- 25 e) LpTK 4 (SEQ ID NO:14);
 - f) LpTK 13 (SEQ ID NO:16);
 - g) HpTK 5 (SEQ ID NO:24);

```
h) bpTK 1 (SEQ ID NO:25);
i) bpTK 2 (SEQ ID NO:26);
j) bpTK 3 (SEQ ID NO:27);
k) bpTK 4 (SEQ ID NO:28);
l) bpTK 5 (SEQ ID NO:29); and
m) bpTK 7 (SEQ ID NO:30).
```

- 4. Isolated DNA of human origin which comprises a DNA fragment whose sequence encodes an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein tyrosine kinases.
 - 5. Isolated DNA of Claim 4 which encodes an amino acid sequence selected from the group consisting of:

```
a) SAL-S1 (SEQ ID NOS:6 and 18);
```

- 15 b) SAL-D4 (SEQ ID NO:8);
 - c) LpTK 2 (SEQ ID NOS:10 and 20);
 - d) LpTK 3 (SEQ ID NO:12);
 - e) LpTk 4 (SEQ ID NO:14);
 - f) LpTK 13 (SEQ ID NO:16);
- g) HpTK 5 (SEQ ID NO:24);
 - h) bpTK 1 (SEQ ID NO:25);
 - i) bpTK 2 (SEQ ID NO:26);
 - j) bpTK 3 (SEQ ID NO:27);
 - k) bpTK 4 (SEQ ID NO:28);
- 25 1) bpTK 5 (SEQ ID NO:29); and
 - m) bpTK 7 (SEQ ID NO:30).

- 6. A homogeneous protein of human origin which includes an amino acid sequence exhibiting sequence homology with the catalytic domain of tyrosine kinases of the c-kit family.
- 5 7. A homogeneous protein of Claim 6 in which the amino acid sequence is selected from the group consisting of:
 - a) SAL-S1 (SEQ ID NOS:6 and 18);
 - b) SAL-D4 (SEQ ID NO:8);
- 10 c) LpTK 2 (SEQ ID NOS:10 and 20);
 - d) LpTK 3 (SEQ ID NO:12);
 - e) LpTk 4 (SEQ ID NO:14);
 - f) LpTK 13 (SEQ ID NO:16);
 - g) HpTK 5 (SEQ ID NO:24);
- 15 h) bpTK 1 (SEQ ID NO:25);
 - i) bpTK 2 (SEQ ID NO:26);
 - j) bpTK 3 (SEQ ID NO:27);
 - k) bpTK 4 (SEQ ID NO:28);
 - 1) bpTK 5 (SEQ ID NO:29); and
- 20 m) bpTK 7 (SEQ ID NO:30).
 - 8. A protein of human megakaryocytic origin which exhibits significant sequence homology with the FLT/FLK family of protein tyrosine kinases.
- 9. A protein of Claim 8 encoded by the nucleotide sequence (SEQ ID NO:5).
 - 10. A protein of Claim 8 encoded by the amino acid sequence (SEQ ID NO:6).

- 11. A protein of Claim 8 encoded by the nucleotide sequence (SEQ ID NO:17).
- 12. A protein of Claim 8 encoded by the amino acid sequence (SEQ ID NO: 18).
- 5 13. A protein of human megakaryocytic origin which exhibits significant sequence homology with the FGF receptor family of protein tyrosine kinases.
 - 14. A protein of Claim 13 encoded by the nucleotide sequence (SEQ ID NO:7).
- 10 15. A protein of Claim 13 encoded by the amino acid sequence (SEQ ID NO:8).
 - 16. A protein of human megakaryocytic or lymphocytic origin which exhibits significant sequence homology with the NGF receptor family of protein tyrosine kinases.
 - 17. A protein of Claim 16 encoded by the nucleotide sequence (SEQ ID NO:11).
 - 18. A protein of Claim 16 encoded by the amino acid sequence (SEQ ID NO:12).

- 19. A DNA expression vector containing a DNA sequence of numan origin which hybridizes to a DNA fragment which hybridizes to DNA encoding an amino acid sequence present in the catalytic domain of a protein tyrosine kinase of the c-kit subgroup of protein kinases.
- 20. The DNA expression vector of Claim 19 containing a DNA sequence selected from the group consisting of:
 - a) SAL-S1 (SEQ ID NOS:5 and 17);
 - b) SAL-D4 (SEQ ID NO:7);
- 10 c) LpTK 2 (SEQ ID NOS:9 and 19);
 - d) LpTK 3 (SEQ ID NO:11);
 - e) LpTk 4 (SEQ ID NOS:13 and 21);
 - f) LpTK 13 (SEQ ID NO:15);
 - g) LpTK 25 (SEQ ID NO: 22); and
- 15 h) HpTK 5 (SEQ ID NO: 23).
 - 21. A cell transformed by the expression vector of Claim 19.

sals1 (160 bases) FLKI-LIKE

PTK1/3 PRIMERS 41 5'ggatcctgtgcatcagtgacttagggactaggaacattctgctgtcggaaagcgacgtggt W S 101 8 æ 81 21 O

gaagatetgtgaettttggeettgeeegggaeatetaeaaagaeeeegetaegteegeaa

61

 \odot

S

۵.

0

gcatgcccggctgccctgaagtggatggcgccagaattc 3 ¥ 121

FIGURE

S

0

bases)sald4 (147 FGFR-LIKE

;

6' ggatccattcacagagacctagcacgcascatcctggtctcagaggacctggtaacc ш × ĸ 21 ~ v

ggcctggccaaagccgagcggaaggggctagactcaagccggctg 101 IJ 8 ш ¥ 81 5 aaggtcagcgacttt **11** > ¥ 61

cccgtcaaatggatggctcccgaattc ш 141 I 121

LpTK2

GTTGGAATTCCTTCCGGCGCCATCCATTTCACCGGCAGCTTTATTTCGTGTCTAGATTCA

TAGATGTCTTCATTATCTACCTTAAAAACTCTGGCAAGTCCAAAATCTGCTACTTTGTAG

ATATTATGTTCACCAACGAGGACATTCCT

FIGURE 3A

LpTK3

GTGCACAGGGATCTCGCGGCTCGGAACATCCTCGTCGGGGAAAACACCCTCTCGAAAGTT
GGGGACTTCGGGGTTAGCCAGGCTTATCAAGGAGGACGTCTACCTCTCCCATGACCACAAT
ATCCCCTACAAATGGATGGCCCCTGAGGGAA

FIGURE 3B

LpTK4

FIGURE 3C

LpTK13

GTCAATCGTGACCTCGCCGCCGAAATGTGTTGCTAGTTACCCAACATTACGCCAAGATC
AGTGATTTCGGACTTTCCAAAGCACTGCGTGCTGATGAAAACTACTACAAGGCCCAGACC
CATGGAAAGTGGCCTGTCAAGTGGTACGCTCCGGAATGCATCAACTACTACAAGTTCTCC
AGCAAAAGCGATGTCTGGTCCTTTGGAATTC

FIGURE 3D

GTTACATAAC CAATGTATTG GCACTTCCCC GCCCATATAT TTAGTTCATA AATCAAGTAT TACCCCCACT AGTAATCAAT TCATTAGTTA AGTTATTANT TCANTANTTA ATTATEGAČT I TAATAACTGA Trecacere ceceaente

_

GGACTTTCCA CCTGAAAGGT ACCCCANTAG TCCCATAGTA AGGGTATCAT FCACCTATGT ACTGCATACA ACCICAATAA CCCCCCATTG CCAACGACCC TTACGGTAAA TGGCCCGCĈT GGCTGACCGC 101

CAATCACGGT GTTACTGCCA CTATTCACGE AGTACCCCCC TCATATGCCA ACTATACGCT ATCAACTGTA TTGGCAGTAC AACTGČCCAC TTGACGGGTG TGGGTGGAGT ATTTACGGTA ACCCACCTCA TAAATGCCAT TTGACGTCAA AACTGCAGTT 201

ATCCTCATCC TACCACTACC CCCTATTACC TATTAGTCAT ATAATCAGTA TACATCTACG TACTTGGCAG CCCTCAAAGG ATCACCTTAT TACTGGAATA AANTGGCCCG CCTGGCATTA TGCCCAGTAC TTTACCGGGC GGACCGTAAT ACGGGTCATG 301

TTTGGCACCA AAACCCTGGT GGGAGTTTGT TCACCTCAAT TCCACCCCAT ACCTGGGGTA TTTCCAACTC AAAOCTTCAG CACACCCCCA GGTTTTGGCA GTACATCAAT GGGCGTGGAT AGCGGTTTGA 401

CAGAGCTČĞÎ **TCTATATAAG** AGATATATTC CGCACCCTCC TAGGCGTGTA AAATGGGCGG TTTACCCGCC CCATTGÁCGC GCTAACTGCG TCCCCC AGGCGG CAACT GACTITCCAA AATGICGIAA CICAAAGGII TIACAGCATI AAATCAACGG (TTTACTTGCC (591

CTTGCCACGT GATCCAGCCT CACCGGGACC CCATAGAAGA CANANCTOCA CATCCACCCT CTGCACACGC TTAGTGAACC GICAGATCCC AATCACTTGG CAGTCTAGGG 601

						-
ACN'FA TGTAT	CACCT	agttggtgga Tcaaccacct	GTCGGCGTAG	CTGGC	AGGCG	TTACE
TFANTACA'FA Aattatgeat	AACTGCACCT TTGACGTGGA	AGFFG	GTCGG	ACGCACTGGC TGCGTGACCG	CAUGCAGGG	ATGTATTACE
ACAA		GATA	CCAG			2446
GCGGCTACAA CGCCGATGTT	TCCCAGGTCC	atcactgata tagtgactat	GTTGAACGAG Caacttgctc	GCGCTGCTCG	CCCCCAGCTT	CTACTCTTTG
	<u> </u>					
tcgttagaac agcaatcitig	AGGTGTCCAC TCCACAGGTG	CCCATTCTTT	CCCTAGACCT	GAACAAGCGG	actgggaatg tgacccttac	CGCGGCCGCA
CCAL TTGGCT GGTGAACCGA	Trctctcac	TGTGCTGGCG	ATCCGTGCCG TAGGCACGGC	GGCACTTCAG CCGTGAAGTC	CTCATTTCTG Gagtaaagac	GCCTCCAGGT
GTCTATAGGC CAGATATCCG	CACTITGCUT	CGAGATCCAT	CAATACAGTG CTTATGTCAC	GCGCTTTACT CGCGAAATGA	ACGACTGGCG	CCAGTTCTGC GCTCAAGACG
CCCCTATAGA	GAATAACATC CTTATTGTAG	CCCTCGACCT	AGTTGCAGCC TCAACGTCGG	TCAGCAGCCG	AGAGCCGACG	TCCATACCTA AGGTATGGAT
0 0 0 0 0 0 0 0 0	CTTA		ACTT	TCAG(AGTC(AGAGO TCTC	
acgtaagtac Tgcattcatg	TGACACTATA ACTGTGATAT	ctctagagat gagatctcta	ACCATGACAA TCCTACTGTT	CCCCA	70000 A0000	CGAGGGATCT CCTCCCTAGA
ACG'FA TGCAT	TGACA	CTCTA	ACCAT TCCTA	GCTTGGGGGT	TTCGGTGCCG AAGCCACGGC	CGAGO
ACTC TCAC	TAGG	SGATC CTAG	GTCA			SATCT
GCCAAGAGTG CGGTTCTCAC	accatttagg tgctaaatcc	CCCCCCTAG	Taaagtgtca Atttcacagt	CTGGCGGAAC	ATCATAGCAC TACTATCGTG	CAATGGATCT GTTACCTAGA
			7			
GATTCCCCGT CTAAGGGGCA	TCATACACAT AGTATGTGTA	ga t tgaattc Ctaacttaag	ttatcagtga aatagtcact	GACACGCAAA Ctgrgcgttt	CTGGCGGAGA	ACCCCCACCA
TTGGAACGCG AACCTTGCGC	accttatgta Tggaatacat	CGG TTCTATC GCCAAGATAG	Catattatgt Ctataataca	ACGGTCTGAC TGCCAGACTG	CGAAGCCATG GCTTCGGTAC	CTGCTCGCCT
		901 CG CC				
701	801	õ	1001	1101	1201.	1301

AAGGGCGTCT CCCTTTCCTT TTCCCGCAGA GGGAAAGGAA

CAGACCATGA

TGTTACTCAG ACAATGAGTC

GCANGGCACC

CCAGTGTCTG

CATANGGGCC GCTTGAGGGC TCTTTGGTCA AGCAGTAACG GTATTCCCGG CGAACTCCCG AGAAACCAGT TCGTCATTGC

1501

TAGCGANAGT

GTAGTAGATA GCAGAAGAAA CATCATCTAT CCTCTTCTTT

TTGGAAGTTT

AGAACCGCAG CACCAAGGAG

GIGCTICAGI GGICACACIC CITCICCACI TICATGCICC CACCCAAGICA CCAGICICAG GAACAGGIGA AAGIACGAGG

1701

Grcacgergg Cagtgcgacc	
GTGACCTAGA AGGCAAGAGG TGAGCCCTCT GTCACGCTUG CACTGGATCT TCCGTTCTCC ACTCGGGAGA CAGTGCGACC	
CHCACCTAGA AGGCAAGAGG TGAGCCCTCT	
CTGAAGGGAC ATTGTGAGAA	
CCCCCCCTCACTC CTGAAGGGAC ATTGTGAGAA	
GGGTCAGGTG	
CCCCCCCACA	
AGGAATAACT	
1401 CATATTACCA AGGAATAACT GTATAATGGT TCCTTATTGA	
1401	

CCTCTC	
CCTC	
CCCTCCCC ACTCATCCGC ACCCTAACC CCTCCCTGTC	
36 AG	
agtcatcc tcagtagg	
ນນຸນ	
)))))))))))))))))))	
TECTCCACCA GCTTCTTGTG GGAGGCTGGA TATTATCCAGACGTGGT CGAAGAACAC CCTCCGACCT ATAATAGGTC	
TATT	
766A ACCT	
CAGGC	
STS SAC SAC	
FTC#7 NAGAA	
200 ≰.t.	
CCACO	
TCCT	
CACTC	
GGGAA	
CTCA	
CCAGCAGTCA GGGAACACTC	
1601 CCAGCAGTCA GGGAACACTC	

ACACACACAC CAAGACCTAC CAGAACCCGC TITICITGICT AGAGACACACAC O R P G S E 0 R	
TTTC AAAG S E	
ပ္ပပ္ပံ ပုပ္ပံပံ	
CTCAAGCCG GACTTCGGC	
55€	
SCTAC CGATC	
TTCA	
27. 20.00	
HCTC NGAG	
SAAG	
CTGC GACGC	
7 Y	
ACGT	
AGAGAAGAAA TGCTGACGTA TGCTGCCTTC	
4. tr	
SAAG	
AGAC	
GTGC	
ttataagtge aatattcaeg	
CTTT	
PTGAT	
1801 CFTAAAGTCF FIGATCTFFC	
AAAG	
£3	_
1801	349

CAACAGGIGT CTCGGAAACA TCCAGCGAACC CCAGTACCCCCAACAGGIGT CTCGGAAACA TCCAGCAACC CCAGTACCCCCAACCC CCAGTACCCCCAACCCCCAACCCCAACCCCAACCCCAACCCCAACCCC	GTAATACČTG GCGGCCAGGC CATTATGGAC CGCCGGTCCG Y Y R A A L
CACCCPPTCT CTCCCAAACA 8 G K	CCGGGAAAGG ACACCCAGTT GGCCCTTTCC TGTGGGTCAA G P F S V W N
TCTCTCTCTG ACAGACAGAC S D T Q	CAGCCCTCT GGCCAGGCAC GTCGGGGAGA CCGGTCCGTG
ACCATCCCAC TGGTAGGGTG V W G	CAGCCCCTCT GTCGGGGAGA R A G R
CACACTCCTC CGAGGCCAGC ACCATCCCAC TGTCTGTCTGGTGGTGAGGGGGGGGGG	CCACGGTCT CAGCCCCTCT GGCCAGGCAC GCTGCCCAGA GTCGGGGAGA CUGGTCCG1G G R T K A G R A L C
CACACTCCTC GTGTGAGGAG E C E E	CCTGGAGGAA GCACCTCCTT R S S
	ATGTCTTCAT TACAGAAGTA F T K H
ATACCTGCTC TCTATCTGCT TATGGACGAG AGATAGACGA Y R S E I O	2001: AAT'FCCTCAA ATGTCTTCAT TTAAGGAGTT TACAGAAGTA
1901	2001: AA TT

TCCACCTCTC ACGTCCAGAG A P R CTCACCTCTG
CACTCGAGAC
E S S 0 CCTCCGAGAA CGAGCCTCTT L E S F CTGCCCTCTT GACGGGAGAA S G B ATCTCCACCA TAGAGGTGGT I E V CTCCCAGAAG CACCCTCTTC 0 S F CACCCCCAGG CACCCCCTCC D G L TCCTCCACAC ACCACCTGTG M T S V CCTGGAGCAG GGACCTCGTC G 0 L L TGTAGGGCCA ACATCCCGGT H L A AGGCCCCTGC TCCGGGACG L G R CTGGGCGATG GACCCGCTAC 0 A I TTCCTCTTGC AAGGAGAACG E E O GGGCTGTCCT CAGCGTCAGC CCCCACAGCA CTCGCAGTCG P S D E A D A ACACCTCCTC ' AGAGCTGCGC GGGGCCATGC TCTCGACGCG CCCCGGTACG S S R P A M TCTGCCCTC CAGGCTTGGC ACACCCCCAC CTCCGAACCC S II R 0 L S P 2201 2/5 2101

TCTCCCTGCT ACACGCACCA S O T T ACCCCCTCCC TCCCCCACCC CACACGTCAC Grerccagte W V D CCCCTCTCTC CGCCAGAGAG G D R CCCNAAGGAC GGGTTTCCTG G F S PCATCCTTGT AGTAGGAACA R H R T AGAGAAGCAC TCTCTTCGTG W L L V TCCGGGGCCC AGGCCCCGGG E P A AAGATCTCCC TTCTAGAGGG AGTGGCCAGC TCACCGGTCG T A L CCCCAGAGAG GGGGTCTCTC G L S GTATGGCGGG CATACCGCCC R I A P ACGGGGAGGC TGCCCCTCCG Y P S A A'FGATGCGGC TAC'FACGCCG M I R ACCCCAGGGT TGGGGTCCCA V G P CCCTTGGGGT CTCCGGACCA GCAGTTCAGG CCCAACCCCA GAGGCCTGGT CGTCAAGTCG 2301 208

ATTGATCTGC TAACTAGACG

AGAACTCCTC
TCTTCAGGAG
C F E E

2401

175

CACCCCCCCCA CACCCCCCT D R A CTTTGTAGAT GAAACATCTA D K Y I TACTCGGGGT ATCAGCCCCA Y D P CTTGCGGACG GAACGCCTGC K R V CCCCTCACTGCC CCCCTGACGG N A S G AGGGGCAGCC TCCCCGTCGG CATCCACTTC GTAGGTGAAG H W K TCCAACATGC TTTCAGGGGC AGCTTCTACG AAAGTCCCCG CATCACCTTG CATGTGGAAC 2501 141 AGGAACTCCA TCCTTGAGGT L P E TCCCCAAGCC ACCCCTTCGG R S A GGATGCACTT CCTACGTGAA H I C K ACCICTCTCT TCCAGAGACA L D R CCCAGCAGCC GGCTCGTCGG GCAGAATGTT CGTCTTACAA L L I N CTTTCCGACA GAAAGGCTGT S E S AGGCCAAAGT CACAGATCTT CACCACGTCG TCCGGTTTCA GTGTCTAGAA GTGGTCCACC 2601 108 CCCGCCTCGC GGGCCCAGCG TCTGGAGAAG ACACCTCTTC D P 3 AGCTTCTTGG TCGAAGAACC A E 0 ACAGGTCCTC TG1CCAGGAG GGGCrCAGCC CCCGAGTCGG P S L CATGGTCAGC GTACCAGTCG H T L MAGATCTTC FFCTAGAAG L D E 55> CTGTAGCAGA GACATCGTCT S Y C 1701 TCCCTCTCGC CACCTGGANG
AGGGAGACCG GTGGACCTTC
15 M G R A V 0 F . .

FIGURE 4E

GCGGAAGCGT CGCCTTCGCA R F R CCACCATGGC GGTGGTACCG B V M A Crescencer Gaccectega R A L CCGATCCAGC GGCTAGGTCG R D L CCGCCCCCT GCCCGCCGA G P R R rccrecrec Accaccace D S S GAGGACCCTG CTCCTGGGAC L V R ACCGCGCGAA TGGCGCGCTT GTCTTCGAGA CAGAAGCTCT T K S TCCCCCCTCG AGGCGGGAGC C G E 2801

CATTCCCCCCC GAGGGGGCTC CAGGGGGGGT CCCCCAGTC **PCCCTCCACC**ACCGACCTCC TCCACGAAGC ACGTGCTTCG CTCCTGCGGA GAGGACGCCT CCCCTCTGAN CCCCTCTGAN CCCCCCTCCT CCCCCCACCA C R O 2901

TTTATTCCAG NANTANCGTC OCCCÁACTO CGGGTTGAAC CCCCCCCTAC **GCAGAAGCTT** CCTCTTCGAA CTCACCTGGA CCCCACTCTA 00000¥00000 CCCCCCCCT CACCGXGCAG CCCCCCCCAC 3001

TCATCAATCT AGTAGTTACA TTGTCCAAAC TAGTTGTGGT CACTGCATTC GCATTTTTT CACAAATAAA GTGTTTATTT TCACAAATTT AGTGTTTAAA AGCAATAGCA TTACAAATAA AATGTTTATT CTTATAATGG 3101,

TCTGAGGCGG AGACTCCGCC TTAGGTACCT AATCCATGGA AGGAACTTGG TCCTTGAACC CCTCTCAAAG CCTCAAATAA AGCACCATGG TCGTGGTACC AATTCGGCGC Traagccgcg ATCGGGAÄTT TAGCCCTTAA GTCTGGATCG CACACCTAGC ATCTTATCAT TAGAATAGTA 3201

TTAGTCAGCA TCCATCTCAA ATCCAAAGCA ACCCACACACA CCTCCCCAGC AACTCCCCAG CCACACCT 10,0 TCTCTCAGTT ACACAGTCAA CTGTGGAATG AAAGAACCAG 1001

ACTCCCCCCA TCACCCCCCCT CCCCCCCCTA CAACCATAGE AATTAGTCAG CATCCATCTC GTATCCAAAG STCCGTCTT ပ္ပပ္ပ AGGCTCCCCA GAAAGTCCCC ACCAGGTGTG TGGTCCACAC 3401

F. K
CC AGTICCGCCC ATICTCCGCC CCATGGCTGA CTAATITITT TTAITTATGC AGAGGCCGAG GCCGCCTCGG CCTCTGAGCT CG TCAAGGCGGG TAAGAGGCGG GGTACCGACT GATTAAAAA AATAAATACG TCTCCGGCTC CGGCGCGC GGAGACTCGA
00 00 00 00 00 00 00
4 33333
CGAG G
NGNGGC PCTCCG
ATGC A
TTA'ITT AATAAA
TTTT VAAAA
CTAAT
CCACT
CCATG
ງວກວກ
ATTCT
99939 222933
AGTTC
22222
AACTO
ACCECECT AACTECECEC A
1501 TCCCCCCCT AACTCCGCCC A
150

AGCTTGGCAC TGGCCGTCGT TITACAACGT CGTGACTGGG TCGAACCGTG ACCGGCAGCA AAATGTTGCA GCACTGACCC	CTANTAGCGA AGAGGCCCGC ACCGATCGCC CTTCCCAACA
MCCCTC ACCCC	NATAGEGA AGAGG
	CTTAATCGCC TTGCAGCACA TCCCCCCTTC GCCACCTGGC GTA
GCTTTTTGG AGGCCTAGGC TTTTGCAAAA AGCTGTTAAC CGAANAAACC TCCCCATCCG AAAACGTTTT TCGACAATTG	Y TCCCCCTTC
AGGCCTAGGC TCCCGATCCG	TTGCAGCACA
GCTTTTT	CTTAATCGCC Gaattagcgg
	4 M
3601 ATTCCAGAAG TAGTGAGGAG TAAGGTUTTC ATUACTUUTC	3701, AAAACCCTGG CGTTACCCA
3601	370]

22	•
G CGGTATTTCA CACCGCATAC GTCAAAGCAA CCATAGTACG CACCATAAAGT GTGGCGTATG CAGTTTCGTT GGTATCATGC C	
GICANAGCAA	
CACCGCATAC	
CCCATANAGE	
CCCATCTGTG	
TTTCTCCTTA AAAGAGGAAT	
EG ANTGGCGCCT GATGCGGTAT TTTCTCCTTA CGCATCTGTG CO	
AATGGCGCCT	Traccord
CTGANTGGCG	SACTTACCCC
1801 CTTGCGTAGC CTGAATGGCG A	CAACGCATCG GACTTACCGC T
3801 G	

9/35 -

CAN
V CO V
13
100 115
33
A CACTIGCCAG CGCCCTAGCG CCCCCTCLTT 1CCCTTCTT T GTGAACGGTC GCGGGATCGC GGCGAGGAA AGCGNAAGAA
30 20 20
ပ်ပွဲ ပြပ်ပွဲ
ၓၓ
25
£25
55 55 55
44
222
25.0
53 88
36
ညည
TA AGCGCGGGGG GTGTGGTGGT TACGCGCAGC GTGACGGCTA CL AT TCGCGCCGCC CACACCACCA ATGCGCGTCG CACTGGCGAT G
100 CC.
55 00
ACA
80
S S S S
000
100
TY.
200 200 200
၁၁၁
1' CGCCCTGTAG CGGCGCATTA I
CT CT
)))))
), (C
3901' CGCCCTGTAG CGGCGCATTA A

OF FOCUCIONE FOUNDATIONS OF TANK CONTROL CONTROL CONTRACTO CTTANCEGON CONTROL
CTTTACGGCA CANATGCCGT
CCANTIAGE
TT TAGGGTTC AAATCCCAAG
00000CTCCC
GCFCTAAATC
TCCCCCTCAA
TCCCCCCCTT ACCCCCCAA
r crescencer
CCCTTCCTTT
4001

CACGTTCTIT AATAGTGGAC GTGCAAGAAA ITATCACCTG CACGGTTTT CCCCCTTTGA CGTTGGAGTC AGFGGGCAT CGCCCTGATA TCACCCGGTA GCGGGACTAT AAAAAACTTG ATTTGGGTGA TGGTTCACGT TTTTTTGAAC TAAACCCACT ACCAAGTGCA 4101

A STATE OF THE PARTY OF THE PAR

	TATFGGTTAA AAAATGAGCT ATAACCAATT TTTTACTCGA	CTGATGCCGC ATAGTTAAGC GACTACGCCG TATCAATTCG	TOTOTGOTCC CGGCATCCGC	ATTCTTGAAG ACGAAAGGGC TAAGAACTTC TGCTTTCCCC	TGFGCGCGGA ACCCCTATTT ACACGCGCCT TGGGGATAAA	AGGAAGÁGTÁ TGAGTATTCA TCCTTCTCAT ACTCATAAGT	TAAAAGATGC TGAAGATCAG ATTTTCTAGG ACTTCTAGTC
	CTAAAGCCGG	ACAATCTGCT TGTTAGACGA	CTGACGGGCT	CGCTCCGTCA	TTCGGGGAAA AAGCCCCTTT	atattgaaa Tataactete	CTGGTGAAAG GACCACTTTC
	GGATTTTGCC CCTAAAACGG	CACTCTCAGT	GACTGCGCGG	ACCGAAACGC TGGCTTTGCG	GGFGGCACTT	TGCTTCAATA	CCCAGAAACG GGGTCTTTGC
	GATTFATAAG C CTAAATATTC C	TTTTATCGTG (CCAACACCCG	CACCGTCATC GTGGCAGTAG	TTAGACGTCA C	CCCTGATAAA	AAAAACCAGT
FIGURE 4G	CTATTCTFFF G Gatargaara C	ACCIPITACAA T TGCAAATGIT A	CCGACACCCG C	CACAGGTTTT	TAATGGTTTC T	GAGACAATAA C	TCCCTTCCTC 1
ţ	CTATUTCGGG C	CANANTAT'IA A Getetatare t	TGGCTGCGCC O	CTCCATCTCT	GTCATGATAA T	ATCCGCTCAT G TAGGCGAGTA C	TCCGCCATIT 1
	ACACTCAACC C TGTGACTTGG G	CCAATTTAAA CGCTTAAAATT G	GACTGGGTCA ⁻ 7	AGAGGCCCTC	ATACCITAAT G TATCCAATTA C	TCANATATGT A	AAGGGAAAAA 1
	AACTGGAACA A TTGACCTTGT 1	AAATTTAAACG C	ATCGCTACGT (TAGCGATGCA)	GCTGTGACCG	CCCTATFTT I	CTAAATACAT 1 GATTTATGTA 1	CAGCGGGAAT
	ACAACAAGGT 1	CTAAATTGTT	CAACTCCGCT	TTACAGACAA AATGYCTGTT	CTCCTGATAC C	CAAATAAAAA C	ACATTTCCGT TGTAAAGGCA
	4201	4301	4401	4501	1091	4701	4801

				11/35			• .
	AGCACTTTT TCGTGAAAA	ttgagtactè aactcatgag	CTTACTTCTG	AATGAAGCCA TTACTTCGGT	CCCGCCAACA	AGCCGGTGAG	ATGGATGAAC
FIGURE 411	TCCAATGATG À AGGTTACTAC 1	AATGACTTGG 1 TTACTGAACC 1	CTCCCCCCAA	ACCGGAGCTG	ACTCTAGCTT	ataaatėteg tatttagace	TCAGGCAACT
	AAGAACGTTT 1 TTCTTGCAAA A	CTATTCTCAG A	AGTGATAACA C TCACTATTGT C	Atcgttggga Tagcaaccct	CGAACTACTT	TTTATTGCTG AAATAACGAC	CCACCCCAC
	TTTCGCCCCG 7	CCCCCATACA C	CATAACCATG 1	ACTCGCCTTG TGAGCGGAAC	TATTAACTGG ATAATTGACC	CCGACCGACC	GTTATCTACA Caatagatgt
	CCFFGAGAGF 1 GGAACTCTCA 2	CAACTCGGTC C	GCAGTGCTGC (CGTCACG)	GGATCATGTA	TTGCGCAAAC AACGCGTTTG	CCCCCTTCC	CCGTATCGTA GCCATAGCAT
	CCCCATICIA C	CGGGCAAGAG Č GCCCGTTCTC	AGAGAATTAT C	ACAACATGGG TGTTGTACCC	GCCAACAACG	CTTCTGCGCT	GTAAGCCCTC
	CATCTCAACA G	GTGATGACGC C	CATGACAGTA A	GCTTTT'FTGC CGAAAAAACG	CAGCAGCAAT GTCGTCGTTA	TGCAGGACCA	GGGCCAGATG
	CATCGAACTG G GTAGCTTGAC C	GTATTATCCC G	TTACGGATGG C	CCTCGATTGG	ACCACCATGC TGGTGCTACG	CCCTATTCA	TGCAGCACTG ACGTCGTGAC
	GACTGGGTTA C CTCACCCAAT G	ATCTCCCCCC C	GAAAAGCATC T	CTCCTGGCTT	CGAGCGTGAC	TCGATCGAGG	GCGCTATCAT
	TTGGGTGGAC G	AAGTTCTGCT A' TTCAAGACGA T	ACCAGTCACA G TGGTCAGTGT C	ACAACGATCG C TGTTGCTAGC (TACCAAACGA C	ATTANTAGAC 1 TAATTATCTG 1	CGTGGGTCTC
	4901 T	5001 A	5101 A	5201 7	5301 7	5401 2	5501 (

11/35

ACAGGAGAGC GCACGAGGG7 TGTCCTCCC

CCGTAAGCGG CAGGGTCGGA GCCATTCGCC GTCCCAGCCT

TICCCGAAGG CAGAAAGGCG GACAGGTATC

TACAGGGIGA GCATTGAGAA AGCGCCACGC ATGTCGCAC'F CGTAACTCTT TCGCGGTGCG

6201

AGCGAACGAC CTACACCGAA CTGAGATACC TCGCTTGCTG GATGTCGCTT GACTCTATGG

TCGGGCTGAA CCGGGGGTTC GTGCACACAG CCCAGCTTGGAACCCAGCCGACTT GCCCCCCAAG CACGTGTGTC GGGTCGAACC

6101 AGACGATAGT TACCGGATAA GGCGCAGCGG TCTGCTATCA ATGGCCTATT CCGCGTCGCC

V	
<u>[1</u>	
\simeq	•
<u> </u>	
-	١
ĹŦ.	

TAAAACTTO	AGACCCCGTA TCTGGGGCAT	TGTTTGCČGG ACAAACGGCC	GCCACCACTT	GTTGGACTCA	
TAGATTGATT T ATCTAACTAA A	ACTGAGGGTC A	ACCCACCAA /	CCGTAGTTAG	GTCTTACCGG	
ATATATACTT : TATATATGAA	AAAAGCAAGG 1	CACCGCTACC	TCTAGTGTAG AGATCACATC	CATAAGTCGT	
AAGTTTACTC TTCAAATGAG	TTAACGTGAG	ACAAAAAAC TGTTTTTTG	ATACTGTCCT TATGACAGGA	TGCCAGTGGC	
CTGTCAGACC GACACTCTGG	CCAAAATCCC	CTGCTTGCAA GACGAACGTT	CAGATACCAA CTCTATGGTT	CAGTGGCTGC GTCACCGACG	
CGTAACCATT	AATCTCÁTGA	GCGTAATCTG GCCATTAGAC	r caccadages	A ATCCTGTTAC T TAGGACAATG	
CACTGATTAA GTGACTAATT	CCTTTTTGAT	TTTTTCTGC	G TAACTGGC'FT	T CGCTCTGCTA A GCGAGACGAT	
ATAGGTGCCT	AGGTGAAGAT TCCACTTCTA	TTGAGATCCT	T TTCCGAAGG	C CTACATACCT G GATGTATGGA	
CTAGCGACTC	AAAAGGATCT	AAGGATCTTC TTCCTAGAAG	I ACCAACTCTT A TGGTTGAGAA	T GTAGCACCGC A CATCGTGGCG	
CTTTATCTGT	TTETTATET	GTTTTCTAGE	1 ATCAAGAGCT TAGTTCTCGA	CTTCTTGAGA	
8603	5701	Toac	5901	6001	

4 J
RE
GU
-
بنا

					TERRESTORY QUEENSTAR AND	ر: زرداسلسهٔ ۱۳ ر ۱۳ رسهٔ	U W J J J J J J J J J J J J J J J J J J	TCACTCATTA	670) GEGAGTEACC TCACTCATEA GGGACCCAG GOTTEACAFF	670
AA CGCAATTAAT TT GCGTTAATTA	N GTGAGCGCAA	AAAGCGGGCA	TCCCGACTGG	ACGACAGGTT TGCTCTAA	TCCAGCTGGC	GATTCATTAA C'faagtaatt	CGCGTTGGUC	CCTCTCCCCG	1 ACGCAAACCG TGCGTTTGGC	6601
NG AGCGCCCAAT	CTTCGCCTTC	AGTGAGGGAG	GCAGCGAGTC	ACGACCGAGC	CCGCAGCCGA	ATACCGCTCG	GACTGAGCTG	TACCGCCTFFT	ATAACCGTAT TATTGGCATA	6501
T CATTCTGTGG	GTTATCCCCT CAATAGGGGA	1CTTTCC1GC AGAAAGGACG	CCTCACATGT	CTGGCCTTTT	TGGCCTTTTG ACCGGAAAAC	TTACGGTTCC AATGCCAAGG	CCCCCCCTTT	ACGCCAGCAA TGCGGTCGTT	CTATCGAAAA GATACCTTTT	6401
ים פכניכניפטענים	GCTCGTCAGG CGAGCAGTCC	TTTTTCTCAT	PGAGGGTCGA ACTCGCAGCT	ACCTCTCACT TGGAGAGTGA	GGGTTTCGCC	TAGTCCTGTC ATCAGGACAG	CCATACATAT	CCTTTGCGGA	6381 GCTTCCAGG	6 3 6 7

CACACACCAA TTGTGAGCGG ATAACAATTT AACACTCGCC TATTGTTAAA TTCTCTCCAA 6701 GIGACTIACC ICACTCATIA GCCACCCCAG (CACCCCAG AGIGACIANI CCGIGGGGTC)

13/35

6801 ACAGCTATGA CCATGATTAC GAATTAA

FIGURE 5A

CAATGIVATTG GITTACATAAC CCTCAAGGCG GGAGTTTCCCC AATICAAGITAT CGGGTATATA CCCCATATAT TLACTTCATA A'T'GCCCCAG'F TACGGGGTCA TUATITACITIA AGTANTICANT AACCTCCACC GGGCTGTAAC TAATAACTCA TCAATAATTA CCCGACATIG AITFAITFGACT ACTITATITAAT TTCGACCTCG

GGACTITICCA ACCCCAATAG TCCCATAGTA TCACCTATGT ACTGCATACA ACGTCAATAA CCCCCCATTG GGCGGGTAAC CCC ACCOGGGGG CCGACTGGCG GGTTTGCTCGG THE COCCUT GOTHINGCOC COANCEA TTACGGTAAA AATGCCATIT 101

CAATGACGGT GTTACTGCCA CTATTICACCIT TCATATGCCA AGTACGCCCC AGTATACGCGG TCATGCGGGG ATCAAGTGTA AACCGICATIG TEGETEGAGT ATTTACGGTA AACTECCEAC ACCEACTE TAAATGCCAT TTGACGGGTG TTGACGTCAA 201

ATGGTGATGC TACCACTACG CCCTATTACC ATAATCAGTA TATTAGTCAT ATCTACATGC TACATCTACG ATGAACCGTC TACTTGGCAG CCCTGAAAGG CCCACTTTCC ACGGGTCATG TACTGGAATA ATCACCTTAT TGCCCAGTAC TTTACCGGG GGACCGTAAT CCTCGCATTA AAATCGCCCC 301

TTTGGCACCA THECAAGTE TECAECECAT TGAEGTEAAT GGGAGTTTGT AAAGGTTEAG AGGTGGGGTA ACTGCAGTTA CECTCAAACA CTCACGGGGA TCGCCAAACT GGTTTTGGCA GTACATCAAT GGGCGTGGAT AGCGGTTTGA CCANANCCGT CATGTAGTTA CCCGCACCTA 401

CAGAGCTCGT GTCTCGAGCA TCTATATAAG AGATATATTC CCCTCCCACC TAGGCGTGTA AAATGGGCGG TTTTACCCGCC CCATTGACGC TITAGITIGG CTGAAAGGIT TTACAGCATT GTIGAGGCGG CAACTCCGCC AATCTCCTAA AAATCAACGG GACTTTCCAA 501

GAACCCTCCA CTTGCCACGT GATCCAGCCT CTAGGTCGGA CACCGGGACC CCATAGAAGA GGTATCTT G GTTTTGACCT CTCGAGACGC CATCCACGCT AATCACTIVG CAGTCTAGCG GACCTCTGCG GTAGGTGCA GTCAGATCGC TTAGTGAACC 601

GCGCCTACAA TTAATACATA CCCCCATGTT AATTATGTAT AGCAATCTTG TCCTTAGAAC GGTGAACCGA CCACTTGGCT GTCTATAGGC CAGATATCCG CGCCTATAGA GCGGATATCT 701 1TYGGAACGCG GATTCCCCGGT GCCAAGAGTG ACGTAAGTAC
AACCTTGCGC CTAAGGGGCA CGGTTCTCAC TGCATTGATG

The second secon

::484.-

CAGTAAACAG

15/35

5B FIGURE

TCCCAGGTCC AACTCCACCT ACGGTCCAGG TTCACGTCCAGG TCCACAGGTG AGGTGTCCAC AAGAGAGGTG TICICICAC GTCMACGGA CACTITICCCT GAATAACATC ACTGFGATAT TCACACTATA TCATACACAT ACGAITTAGG TGCTAAATCC TEGNATACAT AGTATETETA ACCITATGTA 8 O 1

CTTCTTTTTC CCAAAGGGTA TTTTTTGTAGG ANNNANANA CCACTCCACT TTTTTTTTTTTTTTCCTCACTCA ANANANANA CCCTCGACCT GATTGAATTC CCCGGGGATC CTCTAGAGAT CTAACTTAAG GGGCCCCTAG GAGATCTCTA CGGTTCTATC 901

ATGGGCTATT TACCCGATAA CTCAGACTTT ATACCTATTT CATTITICCTA AACTATITITG GTAAAAGGAT 1TGATAAAAC GTTCTCCTCT TTTAITAATT ACTCAGAGT CTAGGCCACA GCAATCTACT AAATAATTAA TGAGTCTTCA GATCCGGTGT CGITAGATGA 1001

TCAGTTCACA AGTCAACTGT ATATACTGAG TATATGACTC TTTTGATTAC AAAACTAATG TTAAGAGATT AATTCTCTAA TAAGAGTTAT ATTCTCAATA AACGTTGGAC TTCCAACCTG SGCM TAGATAATAA CTCATCCGTT TCTGTAAAGA GTGTAAAGGT ATCTATTATT GAGTAG 1101 AGACATTTCT CACATTTCCA

AGAAAAAAAT **AAAATACGTA** TTTTATCCAT ACCTAGTAAA TCCATCATTT TCTCAACAGC AGAGTTGTCG CTGAAGCAAA GACTTCGTTT TYTACTTETT **ANATGAANAA** TCCTTCTCA AGGAAGACGT TYCCATCATT ATGTCTTTT AAGGTAGTAA CATACTGAAG TACAGAAAAA GTATGACTTC 1201

AAAGCACAGG TTTCGTGTCC AAAAATCTC TTTTTTAGAG THINCICITY AAAAGAGAAA TCCTATAAAA AGGATATTT TTGTAGTTAC TTTACCATCA TAGTTCAGTA AATTGTACCG TTAACATGGC TCCAAGTACA ATCAAGTCAT CACGTTAATC AGGTTCATGT GTGCAATTAC 1301

GTCATTTGTC CCACTICITAAT GGTGAGATTA ATTACTTATT TAATGAATAA ATCTGAGGAA GGATTCAGCA CCTAAGTCGT CAGCCTGATG GTCGGACTAC ATTATT TTCATANTAN MGT TCCTGCTGTG CAGCAAAGCA ATCAAATTCC GTCGTTTCGT TAGTTTAAGG AGGACGACAC 1401

TATCTCTTAA ATAGACAATT GAAATGTAAG ACAATAAATG TCTTATTTAC TCTCTTGATC GACATTTCAA CIKITAAAGTT AGAGAAC'FAG ATCTGTTGAT TAGACAACTA 1501 GAAAATGCTA CAACAGTCAC TGAGTAAAAA TTGGACTATC CTFFTACGAT GTTCACTG ACTCATTFTT AACCTGATAG FIGURE 5C

CAATAAATG CGTCACTCCT TTAATATATA GIGGAACTGA CAGGACGTCA CAACGGGTCA GITATTTAC GTTGCCCAGT CTCCTGCAGT CACCITICACT GCAGTGAGGA AATTATATAT TAAAATKIGTC ATTITIOCCAG AAAGAAAAT AACTTGGTTT AGTGTGCTTA TCACACGAAT TITICITITIA TIGAACCAAA 1601

AACATAAAAC TTCTATTTTC GTTGATAACA TCTCCAGTTC TATTTGACTA ACACGICAAC ATAAACTGAT ATATGTCAGG TATACAGTCC TATCAC1TGA ATAG1GAACT TTGAAATAGG AACTTTTATCC CACAAATAAT CTTTTTCATA ATACA16GCC GTGTTTATTA GAAAAAGTAT TATGTAECGG CACAAATAAT CTTTTTCATA 1701

TCTCAGACAA AGAGTCTGTT CTGAATATGA CACTTATACT Ø Ω TTATTTTOCAT AATAAACGTA < Z TCTTATGAAG AGAATACTTC CCAGTGTTCA 0 ATATTCTTCT TATANGANGA 1801 GAATGGATTA TTTGAATTTG TTTTGCTACT TTATTATTTG NAMACGATGA ANTANTAMAC MANCTTANAC CTTACCTAAT 506

TTAACCACAC AATTGCTGTG GATCTTCTAA CTACAACATT TGAGGTTGTA ACTCCAACAT CCATTCCAGC CGTAAGGTCG < GAATCCGAGA CTTAGGCTCT Œ TAGGTCGTTC ATCCAGCAAG TCAAAATAGT CTTCAAGTTT CCAACGCAGT GTCTCAAATG CAGAGTTTAC 4 CCTTCCCTCA 1 **E 3** GAAGTTCAAA × נ Ħ ۵ AGTYTYTATCA 1901 492

TTCATTACTA AAGTAATGAT TAMAACGGTA ATTITICCCAT ပ TGACATTCCG ACTIGITAAGGC C) GACAGTATGG CTGTCATACC Ç Σ Ö TGGACCCGTG ACCTGGGCAC > GTAGACCTAA CATCTCCATT AGTCTATAGT TTTGAGCCAA NAACTCGGTT z TCAGATATCA > **~** ب CACAGTICGA TGCTIGCGGA ACCAACGCCT ۵ 0 <u>-</u> CTGTCAACCT S z U ۲. 459 2001

GTCGAAATAA CAGCTTTATT ACTITICACCGG TGAAGTGGCC > GGCGCAGTCC CCCCCTCAGG < ۵, TATTACTACG ANTGGCTTCG TTACCGAAGC ш < ATAATGATGC ĸ Z ATCCTGAATT TACGACTTAA Ĺ 2101 TTCATAAAGA AGGATTCCAA ATGACCATAC ATCGGACTTA TAGCCTGAAT TACTAGETATO > 3 Ŀ TCCTAAGGTT AAGTATTTCT 425

TCCTCCTGTA ATCITCACCA TACAAGTGGT Ü = TCTACATATT ACATCTATAA Z **NGACGATGAA** TCTGCTACTT > < TTCAGGTTTT AAGTCCAAAA TTTGAGACCG AAACTCTGGC TCTACCTTAA AGATCCAATT **GTCTTCATT'A** CAGAAGTAAT 2 Ш TAACTATCTA ATTCATAGAT S) TCGTGTCTAG AGCACAGATC <u>::</u> 2201 392

TCAGATGGAT AGTCTACCTA ACCTGTTGAG TCCACAACTC O O > CAACCTGTGC CGCCATGTCT GCGGTACAGA ٥ X GTTGGACACG < Q < ATTCCAGAGG TAAGGTCTCC C CAGATAGGCC CTCTATCCGG TTCTCCCAGC CAGATCTCTG TGAATGTAGT TCCGAGACTC AUGCTCTXSAG ACTTACATCA = AAGACCGTCG GTCTAGAGAC ~ z 359 2301

AACACKCATAA THYTHOTATT

CTAAAGTCCA CATTTCACGE

ATTRICATOTT
TANCOTAGAA

TTATTATT

AATAATAAA

TCAACTCTGT

CCATGTCTCA

GTGTCATTTT GGAGATATTC TINCAGACTT CACAGTAAAA CCTCTATAAG AACGTCTGAA

50

FIGURE

GGTACAGAGT

0

þ

Z

ANANCTAGGT

325

TYTTTCATCCA

2401

ACITY:ACIACA

U

ĹĽ

Ω

۵

TCTCCCATTG

TTGCGGTCTA

GCTTCAGAAG CTGTATGGAG

GATCCCAATC CTAGGGTTAG

NACCCCAGAT

GACATACCTC

CGAAGTCTTC

~

S

TTTGACTGGT

TATICAAGCGG ATACTTCCCC

AGACCTTCCC TCTCCAAGGG >

3

C

3

z

z

H

۲

>

259

225

TAACAAGGTG

CACCTCACCA

C'IGGAGTCGT ATTGTTCCAC

2601

AAACTGACCA

0

AGAGGGTAAC

AAGTGACGAT

×

£-

×

O

S

r

Ω

<u>-</u>

z

a

-

=

O

×

=

ے

×

__

292

TYCACTGCTA

TTTTAATGTT AAAATTACAA

TTOAACCTGG

AACTTGGACC

AAACCTAGGT

GTCCTTCAGT

CACCAAGTCA

CICCCICCT

2501 AGCTGGATAA GCTTTGGATG TCTTAGGTTC TTCATTATCT

CACCGAGGGA

AAGT'AAT'AGA

AGAATCCAAG

CGANACCTAC

TUCHACCTAIT

TITICGATCCA

17/35

GAACCACATC CTTGCTGTAG ACAACTGCTC TGTTGACGAG CGTCACTTGT GCAGTGAACA CTAGTCTTTT CATCACAAAA Ω Ö ACACACAGGC Teretreres TTTTAATTCT ANNATTANGA U > CCCCAGCTTG GGGGTCGAAC TCATCCAGTC AGTAGGTCAG Ç TCGTACCAAA AGCATGGTTT × AAATCCCCCT TTTAGGGGGA ۵, U TCGATCTTTA ACCTAGAMAT GCGTGAGAAA CGCACTCTTT ¥ 0 2701 GTCCACGGTT TTATACGACA NATCAAATGG AGCTGGGACC TCGACCCTCG ATTCTTCTTC TAAGAAGAAG > < ۵. TTAGTTTACC ACAACTTTTC ٦ AATATGCTGT TAAGCAAGTC >-CAGGTGCCAA F

× ĸ TAGATACATA ATTECTTOR TOTTCAAAAG J z E. ACCGAGIRGIT 2801 192

CAAAGAGACG GTTTCTCTCC O ATATTATCAA TATAATAGTT CTTTTTCTCAA CAAAAGACTT ω z AACCGGTCTT TINGCCAGAA H Ö ATTACAAAGG TAATCTTTCC ACTITICTOR TGANAGAGAC CATCTAAAAC TGAAAGAGAG AATTCTCCTT TTTGGCTTTC AAACCGAAAG 0 TTAAGAGGAA Ŀ G'FAGATTTING ACTITICACTC > <u>_</u> ပ 2901 159

TTGCTGACTG AACGACTGAC O 0 ACCCTTGTAG TCGGNACATC 0 ပ CTTCCTTATA S P I GAAGGAATAT GTGCATCAAT CACGTAGTTA z > > TCTCCTCAGC ACAGGAGTCG < w ۵ <u>~</u> TCTAGGCTTC ACATCCGAAG S 0 3001 ATCTGATCTT CCGATTGCTC CANAGAACCA CGGCTCTGCC CCCGAGACGG < <u>'-:</u> = GTTTCTTGGT 3 Ŀ GGCTAACGAG < ပ TAGACTAGAA Ξ 125

CTCAAGTCCT GAGTTCAGGA S TGCTCGGAAG ACGAGCCTTC Ľ. < GTTTGTCACC CNAACAGTGG Ö ۵ AGAACTTGAA TCTTGAACTT 0 CAAAGTGTCC GTTTCACAGG AGCCCTCATG TCCCCAGTAC Ö 3 3101 GAGCCATCTC GTCTTTTCTC CAAGTGTCTG GCAAACCACC CCTTTTCCTCC CAGAAAAGAG GTTCACAGAC Œ Œ CTCGGTAGAG 92

FIGURE 5E

S TCACGGTTCA ACTCCCAACT GCATTTTCAA CCTAAAAGTT AAGGCCCCCT TICCCGGGGA Ö GGGGAGAGCA CCCCTCTCGT 0 CTCTCTGACT GAGACACTGA **GTGGCCATGC** CACCGGTACG GGTGTTTCAT CCACAAAGTA TCAAACAAAG TCGCACCATT AGTITGTITC = AGCCTGGTAA >-0 CAGGAGTCCG GTCGTCAGGC 3201

AGNAGACGAG TCTTCTCCTC CTTCCTCCCC GAACCACCCC GAGTAACACG CTCATTGTGC ACAGATGTTG TCTCTACAAC AGAGCCTCTG TCTCGGAGAC 0 3 AGGTACTCCC TCCATGAGGG 3301 CTTGTCTVICC TCCGTGGACA AACAGGGGAG ATAGGGTTCT GAACAGACGG AGGCACCTGT TTGTCCCCTC TATCCCAAGA W 25 CGGAGAGTAT GCCTCTCATA CACCATACTT GTGGTATGNA AGCACCAACT TCGTGGTTGA CTTTCAAGTC GAMACTTCAG CACCAGGCAA GTGGTCCGTT TATCTTCCTT CCACCT CCTCCA CCAT CAGGGCTICT CCCTCTCCCC TTAGTCTCTG GTCCCGAAGA GGGAGAGGGG AATCAGAGAC CAGGGCTFCT 3401

GCTACCCCGA CGATCGGGCT CCGACCTCCA GGCTGGAGGT ACGAAAGACA rochmerer CTTACCGGCT CCTCGAGAGA GCAAGTCCTA GCAAAGTCCC GTTTCAGATC AGTCCAGCAG CTGGGTTGCA CGTTTCAGGG CAAAGTCTAG TCAGGTCGTC GACCCAACGT 3501

AAATTAAAAG TITANTITIC **ANATANATA** TYTYTYTYT CCCACACAA CCCTCTTGTT GCTTAAGAAT CGAATTCTTA CTTCTTATCT GGTGTCTTTT CCACAGAAAA CCCAAAACTG AGCAGGAGCT GGGCAGCTGC TCACTAGGAA 3601

ACCATTTTGG TGGTANAACC TACTAACCCC AAGAGGAAGC TTCTCCTTCG AAAAGTTAGC TGTCCCAATA TCTTGCCTTT GCCTTTATTT AGACAAATAT CTGAGAACAG AATGGTGCCA CCCAAAATAAA TCTGTTTATA GACTCTTGTC TTACCACGGT 3701

GACCTGCAGA CTGGACGTCT CTCTAGAGTC GCGGCCGCGA CCTCCAGGTC CAGITICIGEG CCATACCTAC KATCTT CTCCCTAGAA CYC TCCACGTCTT GCTTTCGCCA GGGTCGACTC 3801

ATAAAGCATT TTTTTCACTG AAAAAGTGAC TATTTCGTAA AATTTCACAA ATCGTAGTGT TAGCATCACA AATAAAGCAA TTATTTCGTT AATGGTTACA TTACCAATGT AGCTTGGCCG CCATGGCCCA ACTTGTTTAT TGCAGCTTAT TCGAACGAATA ACGTCGAATA 3901

ACCTCTGAAA TGGAGACTTT

GCCTGAAATA

CAGCACCATG

GATCGGGAAT TAATTCGGCG CTAGCCCTTA ATTAAGCCGC

ATCATGTCTG

TATCTT

NATY

GTANGATCAA CACCAAACAG GTTTGAGTAG

CATTCTAGIT GTGGTTTGTC CAAACTCATC

4001

TTACATAGAA

FIGURE

TAGTACAGAC

CAGGCAGAAG

GGCTCCCCAG

AAAGTCCCCA

GTGTGTCAGT TAGGGTGTGG CACACAGTCA ATCCCACACC

GCTGTGGAAT

GAAAGAACCA

AAGACTCCGC

GAGGAACTIG GITAGGIACC ITCTCAGGCG CICCTIGAAC CAATCCATGG AAGACTCCGC

4101

CAATTAGTCA

GCATGCATCT CGTACGA (

AGTATGCAAA TCATACGTTT

CAGGCTCCCC AGCAGGCAGA GTCCGAGGGG TCGTCCGTCT

GGAAAGTCCC

NACCAGGTGT TTGGTCCACA

TATGCAAAGC ATGCATCTCA ATTAGTCAGC ATACGTTTCG TACGTAGAGT TAATCAGTCG

TATGCAAAGC

4201

GTTAATCAGT

TTTATTTATG AAATAAATAC

ACTAATTTTT TGATTAAAAA

CCCATGGCTG

CATTCTCCGC

CAGTTCCCCC

TAACTCCGCC ATTGAGGCGG

ATCCCCCCC TAGGGCGGGG

AGGCGGGGA TTGAGGCGGG

CGTTGGTATC

4301

GCAACCATAG TCCCGCCCT AACTCCGCCC

19/35

CGTAATAGCG GCATTATCGC GTCGAACCGT AAGCTCTTAA GCGCTCGACC CGCCAGCTGG ATCCCCCTTT TAGGGGGGAA CTTTTGCANA GAGGCCTAGG CTCCGGATCC ACTTAATCGC CTTGCAGCAC TGAATTAGCG GAACGTCGTG CCGAAAAAAC GTAGTGAGGA GCGTTACCCA CGCAATGGGT GAVAACCCTG CTTTTCGGAC GGTCTT ATAA CINGCCGTCG TITITACAACG ICGIGACINGG AGCACTGACC GTCTCCGGCT CCGGCGGAGC CGGAGACTCG

GCCTTTTTG

TATTCCAGAA

CAGAGGCCGA GGCCGCCTCG GCCTCTGAGC

4401

GACCGGCAGC ANANTGTTGC

4501

GCCCTATTTC CCCCATANG ACCCATCTGT MAAGAGGAA GAATGGCGCC TGATGCGGTA CTTACCGCGG ACTACGCCAT GGACTTACCG CCTGAATGGC AGTTGCGCAG TICTCCGGGC GTGGCTAGCG CGAAGGGTTG TCAACGCGTC AAGAGGCCCG CACCGATCGC CCTTCCCCAAC 4601

ACACTTGCCA TGTCAACGGT GOTGTGGTGG TTACGCGCAG CGTGACCGCT GCACTGGCGA AATCCCCCTC CCACACCACC AAGCGCGGCG GCGCCCCATT CCCCCCCTAA ACCATAGING GCGCCCTGIA TGGIATCATG CGCGGGACAT ACACCGCATA CGTCAAAGCA ACCATAGTAC TGTGGCGTAT GCAGTTTCGT 4701

And the second s

FIGURE 5G

GAAATCCCAA CTTTACGGTT CCCCCCGAGG AGCTCTAAAT TCGAGATTTA TYCCCCGTCA AAGGGGCAGT AAGCGGCCGA TYCCCCCT TCTCCCCACG AGACCCCTCC AGGGAAGGAA TCCCTTCCTT MGCGMANGA GCCCGCTCCT TTCGCTTTCT CGCGGGATCG CGGGGGAGGA GCCCCTACC 4801

TCCCCCTTTC AGCGGGAAAC AGACGGTTTT TCTGCCAAAA TCCCCCTCAT AGCGGGACTA TAGTCGCCCA ATCACCCGGT ATCCTTCACG TACCAAGTGC CATTTCCCTC CTANACCCAC CAAAAAAACTT GTTTTTGAA TGCAGCTGGG CCGATITIAGT GCITTACGGC ACCTCGACCC GCCTAAATCA CGAAATGCCG 4901

CCCTANAACG TGATTTATAA ACTABATATT CGATAAGAAA GCTATTCTTT CCTATCTCGG AACACTCAAC AAACTGGAAC CTCTTGTTCC TECAACCTCA GGTGCAAGAA ATTATCACCT TAATAGTCCA CCACCTTCTT ACCTTCCACT 5001

CGTGAGAGTC GCACTCTCAG TAAAATACCA ATTTTATCCT TTGCAAATGT **AACCITITACA** ACAAAATATT GCGAATTITTA CGCTTAAAAT AAAATTTAAC TTTTAAATTG TCATTTAACA ACTAAATTGT CTATTGGTTA AAAAATGAGC GATAACCAAT TTTTTACTCG CGATTTCGGC (CCTAAAGCCG 5101

ATCCCCTTAC TAGCCGAATG TGCTCCCGC ACGAGGCCG COCCUMENC GCCCGAACAG GCGCGGGACT CGCGCCCTGA CACCCGCCAA CACCCGCTGA GTGGGCGGTT GTGGGCGACT TCGGGCT CCAGCCCCGA 5201

GCGGATANA CCTCGTGATA CCACCACTAT GACGAAAGGG AAACGCCCCA TTTGCCCCCT GTCATCACCG CAGTAGTGCC GGTTTTCACC CACAGTCT ATCTCTCAGA 7 AGACAAGCTG TGACCGTCTC CGGGAGCTGC TCTGTTCGAC ACTGGCAGAG GCCCTCGACG 5301

TCTAAATACA TGTTTATTTT ACANATAAN AACCCCTATT TTCCCCATAA ATCTCCCCC TACACGCGCC MAGCCCCTT TFTCGGGGAA ACCTCCCACT TCCACCGTGA CTTAGACGTC GAATCTGCAG ATAATGGTTT TATTACCAAA ACAGTACTAT TCTCATGATA TATAGGTTAA ATATCCAATT 5401

TCTCCCCTT ACAGGGGGAA AACATTTCCG ATGAGTATTC TACTCATAAG AAGGAAGAGT AATATTGAAA TTATAACTTT ATGCTTCAAT TACGAAGTTA ACCCTGATAA TCGCACTATT TCACACATA ATAGGCGAGT ACTCTGTTAT TATCCCCTCA TTCAAATATG AAGTTTATAC 5501

CGAGTCGGTT GCTCACCCAA GTTGGGTGCA CTGAAGATCA GTAAAAGATG CATTTTCTAC CGACCACTIT CCTCCTCAAA ACCCAGAAAC **S**H FIGURE TTGCGGCATT TTGCCTTCCT GTFFFFGCTC AACGCGGTAA AACGGAAGGA CAAAAACGAG ATTCCCTTTT TAAGGGAAAA 5601

ATACACCGCG TATGTGGCGC TTTCAAGACG **AMGTTCTGC** CTCGTGAAAA DAGCACTITIT AAGGTTACTA TTCCAATGAT CTTCTTGCAA CAAGAACGTT AAAAGCGGGG TTTTCGCCCC TCCTTCAGAG CCTAGAGTTG TCGCCATTCT AGGAACTCTC AGCGCTAAGA GGATCTCAAC TKITAGCITKIA ACATCGAACT 5701

AGANANGCAT TCTTTTCGTA GTGGTCAGTG CACCAGTCAC CAACTCATGA GTTCAGTACT CTTACTGAAC GAATGACTTG ACTATTCTCA TCATAAGAGT CCCCCCATAC GCAACTCGGT CGTTGAGCCA CGTATTGACG CCGGGCAAGA GCATAACTGC GGCCCGTTCT GGTATTATCC 5801

GGAGGACCGA CCTCCTGGCT ACTTACTICT GACAACGATC TGAATGAAGA CTGTTGCTAG ACTGCGGCCA TGACGCCGGT GAGTGATAAC GGTATTGGTA CCATAACCAT TCCAGTCCTG CGTACTGTCA TTCTCTTAAT ACGTCACGAC **AAGAGAATTA** GCATGACAGT CTTACGGATO 5901

ACGAGCGTGA TGCTCGCACT ATACCAAACG GAATGAAGCC AACCGGAGCT CATCCTTCCC CTAGCAACCC AACTCGCCTT TTGAGCGGAA AGAGCTAAC CGCTTTTTG CACAACATCG GGGATCATGT TAGTACA ည TCCTCGATIG GCGAAAAAC GIGIIGIACC 6001

CTGGATGGAG GACCTACCTC AATTAATAGA TTAATTATCT TCCCGGCAAC ACCCCCTTC TACTCTAGCT ATGAGATCGA CCGAACTACT CCCTTCATCA CTATTAACTG GTTGCCCAAA CAACGCGTTT CACCACGATG CCTGTAGCAA TGGCAACAAC GTGGTGCTAC GGACATCGTT ACCGTTGTTG 6101

CCCCCTATCA GCCCCATAGT CCCACCCAGA CCCTCCCTCT GTTTATTGCT GATAAATCTG GAGCCGGTGA CTCGGCCACT CTATTTAGAC CAAATAACGA CGCCTGCCTG CCCCACCGAC TCGGCCCTTC CCCCAAAG ACC ACTITCTGCGC TGAAGACGCG AACGTCCTGG GCGGATAAAG TYGCAGGACC CGCCTATTIC 6201

TCTAGCGACT AGATCCCTCA CGAAATAGAC ATACCTACTT TATCCATCAA CAGTCCGTTG GTCAGGCAAC ACCACCCCCA TGCTGCCCCT AGTTATCTAC 6301 TIKICAGCACT GGGGCCAGAT GGTAAGCCCT CCCGTATCGT AACGTCGTGA CCCCGGTCTA CCATTCGGGA GGGCATAGCA

! | i

ATTTTCCTAG TAAAAATTAA ATTITION AATTTTCAAG TTAMACTIC TTAGATTCAT AATCTAACTA CATATATACT GTATATATCA ACTIGACACA CAAGATTACT GTTCAAATGA TGACAGTCTG CTATCCACGG AGTGACTAAT TCGTAACCAT AGCATTCGTA TCACTGATTA 6401 CATAGGTCCC

TTTCCTAGAA **NAGGATCTT** TCTTTTCTAG AGANAAGATC GTCTGGGGCA CAGACCCCCT CACTGAGCGT CANAAGCAAG CTTTTCGTTC CTTAACGTGA GAATTGCACT ACCANANTCC TAGGTGAAGA TCCTTTTTTGA TAÅTCTCATGATCCACTTCT AGGAAAAACT ATTAGAGTAC 6501

ATGGTTGAGA TACCAACTCT GATCAAGAGC TTGTTTGCCG GTCGCCACCA AACAAACGGC CAGCGGTGGT CCACCCCTAC TIGITITIT AACAAAAAA GCTGCTTGCA CTTGAGATCC TITITITITICE CGCGTAATCT GAACTCTAGG AAAAAAAGAC GCGCATTAGA 6601

TCTAGCACCG ACATCGTGGC TCAAGAACTC AGTTCTTGAG GGCCACCACT CCCCTCCTCA GCCGTAGTTA CGGCATCAAT AAGATCACAT TTATGACAAG AATACTGTTC CATTGACCGA AGTCGTCTCG CGTCTATGGT GCAGATACCA TITTCCGAAG GTAACTGGCT TCAGCAGAGC AAAAGGCTTC 6701

TTACCCGCATA AATGGCCTAT OCTTODACTC AAGACGATAG CCAACCTGAG TTCTKXCTATC TGTCTTACCG ACAGAATGGC CGATAAGTCG GCTATTCAGC CTGCCAGTGG GACGGTCACC CCAGTGGCTG GGTCACCGAC TCGCTCTGCT AATCCTGTTA
AGCGAGACGA TTAGGACAAT CCTACATACC GGATGTATGG 6801

AGCTATGAGA TCGATACTCT CTACAGCGTG GATCTCCCAC ACTGAGATAC TGACTCTATG CCTACACCGA GCATGTGGCT GAGCGAACGA CCCCAGCTTG CGGGTCGAAC GTCGGGCTGA ACGGGGGTT CGTGCACACA CAGCCCGACT TGCCCCCCAA GCACGTGTGT AGGCGCAGCG TCCCCGTCGC 6901

GGGAAACGCC CCCTTTGCCG AGCTTCCAGG TCGAAGGTCC CGCACGAGGG GCGTGCTCCC AACAGGAGAG TYCTCCTCTC CCAGGGTCGG CGTCCCAGCC GGACAGGTAT CCGGTAAGCG GCCCATTCCC CTCCATA בי AAGCGCCACG CTTCCCGAAG GGAGAAAGGC GAAGGCTTC CCTCTTTCCG Tregegerge 7001

AACCCCACCA TTGCCGGTCGT CCTATKGAAA CCATACCTTT GGGGGCGGAG CCCCCCCTC TCCTCCTCAG ACGAGCAGTC ATTITITICITON TTCAGCGTCG AACTCGCAGC CACCTCTGAC GTCGACACTG TYGTATCTIT ATAGICCTGT CGGGTTTCGC ACCATAGAAA TATCAGGACA GCCCAAAGCG 7101

23/35

FIGURE 53

TTACCGCCTT AATGGCGGAA CTATTCCCAT TGATTCTGTG ACTAAGACAC CGTTATCCCC GCAATAGGGG MGAMAGGAC 7201 ACCCGCCTT TTTACGGTTC CTGGCCTTTT GCTGGCCTTT TGCTCACATG TGCGCGAAAA CGACCGGAAAA ACGAGTGTAC

CGGAGAGGGG TACGCAAACC GCCTCTCCCCC ATGCGTTTGG CGGAGAGGGG CAGTGAGCGA GGAAGCGGAA GAGCGCCCAA GTCACTCGCT CCTTCGCCTT CTCGCGGGTT TUNGTGAGCT GATACCGCTC GCCGCAGCCG AACGACCGAG CGCAGCGAGT ACTCACTCGA CTATGGCGAG CGGCGTCGGC TTGCTCGCTC GCGTCGCTCA 7301

ACCCAATTAA TGTGAGTTAG CTCACTCATT TGCGTTAATT ACACTCAATC GAGTGAGTAA AGTGAGCGCA TCACTCGCGT GAAAGCGGGC AAGGGCTGAC TTCCCGACTG 7401 GCGCGTTGCC CGATTCATTA ATGCAGCTCG CACGACAGGT CCCGCAACCG GCTAAGTAAT TACGTCGACC GTGCTGTCCA ACATGATTAC TGTACTAATG ATTIGTGAGCG GATAACAATT TCACACAGGA AACAGCTATG
TAACACTCGC CTATTGTTAA AGTGTGTCCT TTGTCGATAC 7501 AGGCACCCCA GGCTTTACAC TTTATGCTTC CGGCTCGTAT GTTGTGTGGA TCCGTGGGGT CCGAAATGTG AAATACGAAG GCCGAGCATA CAACACCT

7601 GAATTAA CTTAATT

CAAAGAAGTC CTGAAACTTT GTTTCTTCAG GACTTTGAAA	AGTGGCACAA AACCACACAA TCACCGTGTT TTGGTGTGTT	GAGTTTTGGG GTGCTGCTGT CTCAAAACCC CACGACGACA	ACACTGCCAT CCATCCACCT	
ACCACAAGTA TGGTGTTCAT	CTGGCCCGAG	GTGACGTCTG	r TAACAAGCTC A ATTGTTCGAG	
TAGCAGAAAG ATCGTCTTTC	CGACTTTGGC	TCTAAGGGCA	G GCGTAGCTGT C CGCATCGACA	
AGCGGGGAGG	TGAAGATCAC	CTCCACCTTC GAGGTGCAAG	GTGGCCTAT	
CAGAGCCAGG GTCTCGGTCC	CACAAGACCC CTGTTCTGGG	TTATCAAGGC	A CTGCCTTGCT T GACGGAACGA	
TAGCAGCTGG ATCGTCGACC	CGACATGGAG	GCTCCTGAGG	C GTGGCATTGA G CACCGTAACT	
GGATGGGGĊT	TTGAGAGTGA	A CXCCTGGATG	G XTGCCATACC	
AGAAAGCAGA	CTGCAGCCCA	s cxGGCACCTA	T GACCGGGGAG A CTGGCCCCTC	
CCCCCCCCAC	GGTTTTGCTG	1 ATGAGTGCCG TACTCACGGC	1 GGGAACTGCT CCCTTGACGA	1 6600
-	101	201	301	401

TCAACATUTG CCTGTGATCA AGTTCTAGAG GGAGACTAGT ATATTTGGGA CTATTAGAAA TATAAAGGGT GATAATGTTT 7717766AA46 TOTTHUTTER AUTOACAAAA CONCORDING CACOUNTION ATGACACCET TORCCCCCCA

GEACACGEAA GANGARCTUR AGAATCCCCG TCTTAGGGGC CCATIGGTATE TCATCATATE CCCCCTTCACT : 5 1=1

TGGTCGATGC CTGCAAGTGC TTCCATCACA ATCTATCTCC TACATACACC CACCTTCACC AGGTGTTGTT TGGAGGGGGA 201 GAGAGGGAG AGGTGAGGA CAGTGTAGGA CTCTGGGGTC TCGAGTCCCT GTCACATGCT

CCCAUGGAAC

TTGAAAATGA CATCCACTA CTCAACAAAG CAAAACAGAG TTTTTCATTTTA TETEGGETETT TAAGGACAGE TEGETGAATT AGACCEAGAA ATTEGTETEG AGGGACTTAA ATTTECTTOTE TAAAGGAGAGAG 100

TUCTTACAC ACGAAAAACTG AGAAATACCC TCFFTATGCC ACACTCATTATA TATTGTTTAC AATTACACAA TTAATGTGTT TGAAGCTAFG ACTTGGATGG AGCTEGAGAA TACCTACTTT TTATTGAGAG TCGACCTGTT ATGGATGAAA AATAAGTGTG CALIAAAGUCA 101

GCTTTGCGAT CAGGGGATCC TGGAATGGGT CTGGCCTAGG ACCTTACCCA GAGCGTTCCA CGTATATCTCA AAACCAGGAC GCCCTGCTCT TTTGCTCCTG CGGGACCAGA ATTACAAGA CCTTACTTTA GAAAAATGGA TAATTCT GGAATGAAAT CTTTTTACCT

TGCTGTGCCA GGACATANGG TATTTGGGAC CTTCATGAAT CCTTTTTCAC TTAAAAAGGA CCAGCTCTTC TCACAGGGG AAAGCTGTAA AGAAGAAAGT AGTGGGGG TFTGGACATT TGTTGTTTGA 1 = 3

TAGGGGAAGG TTTCTTAAAG AAAGAATTTC CCCACANTA CCCCCCTCTTAAT AGACCACATT TCTCCTCTAA CAAACTCCTC AGATCTAAAT TCTAGATTTA CHANATORACT GGGGAGGGAA TGGAGGAGGG TGTTGAGAAT ==

: :

:

CTTTCAGATO AGGGCAACTA TCCCGTTGAT GCACTCGAGG AGAAAACAAA TCTTTTCFFT CCTCGGAATT TTCCCCCCTCA CHIMICIATA AGGIGGAAAG CITCITTCATGIT GAAGGAYGGA GAATAACTAACTA TGCAGGTITG GAGAAGTAGA GITGGTACTT 100

TCCTCTTCAA CTACACTTGT ACACCCCATA TGTCCCCTAT CCAACAAAGG ATCATCAGTG TTCCTTTTCT ACTRICATAN CTICATACNE ANCIATICATA CCCTANCACA

ATGAAGAGTT ATTGACCAAT TAACTGGTTA AGATTATGAA TCTAATACTT ATTCAAGTCA TAACTTCACT AATGGTAGGA TTAGGATGGT CCCTAAATAT 1001 ACCATCCCAG TCAATCACCT TTGGTTACCA TCGTAGAAAA TCGTAGGAAAA TCGTAGGACT ACCATCTTT TCATAACGGA AAAAGGGTCT TTTTCCCAGA CCTTGTGAGG AAAATCATTF TTFTAGTAAA CCTTCTCTCG TGTAGGTGGA AGATGCAGGT 1 1 1 1

TTCACCCTGT AAGTCCGACA TACCAAAATG ATGCCCAATT TACGGGTTAA GAAAATGATG ATTCCATGCA TAAGGTACGT GAGAATATAT CTCTTATATA TACAGGATAT CCAAGTTTTG CAATCATAAG CAGGAGGAGAAAG ATGGGTATA GGTTCAAAAG GTTAGTATTC GTGGTCGGTC CCTTCTTCAC TCTTGGACCT CCCATTACCA CCCATCCATA TCCTCTTTCT AAGTGAGGGG ATATAAGAAG GAAACCTCAA GTGCTCGCAG AAGCTTGGGG TATATTGTTC CTTTGGAGTT CAGGAGCGTG TTGGAAGCGG 1001

GAGCAGTACT AGTUCUTUTE TCACCCACAG CACAAACCTG TAACAGAAAA ATAGAAAGGG TATGTTTGCG CCTCAGACCT TCTCCAGGCC CCTTTTAAAC GGAAAATTTG GTGAGAGGAT GGCACATCTT CAATTCCCTT GCTGTGGATA 1501 CTANACATCA CTGAAGCCAT AAAAGCCTTC CTCGTCAAGT CACCAGGTCA R 7C

CTAATTTGTC 11TAACCCTG AAATTGGGAC TCATTGTCCT TUTCTUCTUT AATTGGTGTT TTAAGGAGAA CCTTCCCTT CATCCAAGAC AACATCTCAT TCTATCCAAC 1601

GTGAGGGGAT GGGGATAA TGAGTACTTC-TAGGTTGATT, TCAGAGAATA-TGAATATGAT GAGTGGGGAGTGTATT ACTCATGAAG ATGGAAGTAA AGTCTCTTAT ACTTATACTA AAACCAATTT AGGTATGAAA GCCAGCTACA GAYGGTACAG TTTCGTTAAA TCCAFACTTT CGGYGGATGT GTACCATGT 1061

TITGGANANG TGATGANCGC AACAGCTTAT GGAATTAGCA AAACCTTTTC ACTACTTGCG TTGTCGAATA CCTTAATCGT ATCAGGTGCT TAGTCCACGA AGGTAC'FAGG TCCATGATCC < F INDI GICAAAIGGG-AGITTCCAAG AGAAAATTIA,GAGITTGGG GAACTCAAGA"TGATGACCCA. AAAGAGAGG ACTCATGTCA TTTCTCTCCG TGAGTACAGT GACAGCTCTG 1901,AAACAGGAGT_CTCAATCCAG GTTACCGTCA;AAATGCTGAA;AGAAAAAGCA TTTGTCTCTCA GAGTTAGGTC CAATGCCAGT TTTACGACTT TCTTTTCGT TTTGAATACT GTTGCTATGG TGATCTTCTG **TTACTTGATT** CAGGACCAAT TCCACACTGT O 2001.4GCTGGGAAGCTCACGAGAATA.TTGTGAACCTTGCTGGGGGGG CACAATTTCA GITTTTACCC CACTTTCCAA TCACATCCAA GIGITAAAAGI CAAAAAAAGGG GIGAAAGGIT AGIGIAAGGTT TTTCAAGGAA CCTCTCTCTA PIUL AACTATCTAA GAAGTAAAA AGAAÁAATTT' CACAGGACTT' TICATACATT CTICATTIC TCTTTTTAAA GIGICCIGAA

ATCAAATTGA TACTTTAACT CACTCTGAAG CTTAAGTAAA GGCTTCATGG CCGAAGTACC CAAATCTCAG GTTTAGAGTC CCTGACCCTA 2227 2201/ATTCCAGCAT-GCCTGGTTCA AGAGAAGFFG.AGATACA 17AAGGTCGTA CGGACCAAGT TCTCTTCAAG TCTATGT AGGAATGGAA AAGTTGCCAA TTTGCATATC AAACGTATAG ANTGTECTTA CATTTEANGA TETTETTEC TIACACGAAT GTAAACTTET AGAAGAAACG JIHI ATATGAAAAG CAAAAAGGC TGGAAGAAGA GGAGGACTTG TATAGTTTTG GTTTTTCGG ACCTTCTTGT CCTGCTGAAG

ATGTGACTIT GGATTGGCTC TACACTGAAN CCTAACCGAG TGGTGAAGAT CACCCCATTC GCTTGTCACC CCAGGAACGT 2401-TTTCTGGAAT" TTAAGTGGTG TGTTCAEAGA GACCTGGCGG

TCTACACCAT TTTOAAGGCA AAACTTCCGT CGAAAGCCTG CCTGTAAAAT GGATGGCCCC GGACATTTTA, CCTACCGGGG 2501 GAGATATCAT GAGTGATTCC AACTATGTTG TCAGGGGCAA, TCCCCGTCTG
CTCTATACTA CTCACTAAGG"TTGATACAAG WARACCCGTT ACGGGCAGAC

TTGATGCTAN CTTCTACAAA AACTACGAFF GAAGATGTTT GGCATTCCGG TCCTTACCCT TTGGTGTGAA

7361

2001

TTTTGACTCA AGGAAACGGC AAAACTGAGT TCCTTTGCCG AATGTCCTCA TTACAGGAGT CGTGTTTCGG TGTATCAGAA TGTGGATGGC ACATAGTCTT ACACCTACCG CCTGCTGGGC ATAATGCAAT TATTACGTTA CATCATCGGA AATATACATT TTATATCTAA CTACAGAAGA CARCAGET; GGCAGATGCA CTACAGTCGA CCGTCTACGT CIGATICAAA- ATGGATITAA AATGGATGAG, CCATTTTATG GACTAAGITT TACCTAAATT TIACCTAGIC GGIAAAATA CATCCTTCCC TAATTTGACT TCGTTTTTAG GTAGGAAGGG ATTAAACTGA AGCAAAAATC TTTAGTTTTA AGAGGAACAA TCTCCTTGTT GCCTCAGGTC.GAAGATTCGT CCGAGTCCAG CTTCTAAGCA TCTCTCCGCA ATGGAT, TTGGGGCTAC CTTTCAGCAG:AGAG CACCTACCAA, AACAGGCGAC 1067

CTGCTTCACC ACANAATCTA TTATCAACTG CATCACTAAA AGGTTAATTT TCCAATTAAA TACCABACA JOOL AGGACTTCAT CCCTCCACCT ATCCCTAACA GGCTGTAGAT TAGGGATTGT CCGACATCTA

NIBL, AGACTTTTCT, PCTAGAGAGGG STCTGAAAAGA: PATCTCTCGG

FIGURE 8A

ACCTCGAGGC **GCCGCCCCCA** CCCCCCCCCT ACCCGAGTGA TGGGCTCACT CHARGECCE CCAAACTEAG TTCAGATECT (HTCCCCGGGG (HTTTGAGTC AAGCCTAGGA ACCCCCACCACCCCACC CACACACA CCTCACCCCCC

ATTCCCTCAG TAAGGGAGTC CCACTAGACT TCACCCACTG
A D L K W V T ATTEGANACT
TAACCTTTEA CACACCTCC TCAACACAAA CTCTTYGGACG ACTTGTGTTT E T I. I. N T K 101 GGTGCTGCTC TGCTGGGCTT CGTTGGCGGG AGCTTTGGAA CCAAGCGGGG ACGACGCGAA GCAAGCGGGG TGGAAACGTT < z U >

COCCAGGCCC CCCCTCCGGG CCCACCCCCC GTGACGTGCA CACTGCACGT 0 TACGAAGTGT (ATGCTTCACA) AACAGCACAG CGTGCCGACC TTCTCGTCTC GCACGCGTCG > 0 201 GTGGACGCCC AGTGCGAGGA ACTGAGGGGC CTGGATGAGG CACCTGCCCG TCACCCTCCT TGACTCGCCG GACCTACTCC د ш I 0 >

רדכסססכדה GAGCCCGACC GACAGGGACG L S L P CHOTOCCTOC CGAAGTCGTA CGAGCTCACG CCTCCACTCC U CCTTCACCAT I CCACGTGTAC GCCACGCTGC = CCCTCCCACG < TGACCGAAGC GTGTCCAACC CAGGGTTGCGG CCCCGGCGCACATG 101 ACTEGETTES CACAGGTTES STECCACGGC GGGGCGCGCT 7.2

CTACATCAAG CATCTACTTC ACCICATIOGG CCACCCTOGA TOGAGAACCC GGTCCGCACCT ACCTCTTGGG z ANCOATGOGG ACACGGCCAC CACCCTCACG TUGCTACGCC TGTGCCGGTG CCGGGAGTGC H TUBCTACGCC CCCGACGACG TTCCTCTCGCCCTCTT CTACTATAGGCCGACGACGACG TTCCTCTCGA AGTGCCAGAA CATCATACTC 401 GCGCTCCTGC AAGGAGACCT TCACCCTCTT 105

CCCCACTCCT CCCCTCAGCA GCCCACCGGG AAGGTCAATG TCAAGACGCT GCGTCTGGGA COCAGACCCT O ACTICTOCCA TTCCACTTAC CCCCTCCCCC D CTGGGGGCGA SUI GEGRAVACGG TOGCCGCGGA GCATCTCACC CGGAAGCGCC CACCTGAGGCGCC ACCGCCGCCT CGTAGAGTGG GCCTTCGCGG CGTACAGTCG ACCCCCCCT CACCTUITUSC > 1); <

CAAAAAGTGC GCCCAGCTGA CTGTGAACCT GTTTTTCACG CCGGTCGACT GACACTTGGA 0 GTTTTTCACG CTATCCCTCC ACCTCTTCTA TYSCAGAAGAT GATAGGGACG CATCACCCTG GTACCGGGAC 601 ACCCTVICCTT CTACCTCCC TTCCACCACC AGCCTCCTC TCCCACCCAC AACCTCCTGG TCCCACCCAC AAGGTCCTGG F 0 D 0 0 TUCCAUCCAA GATGGACCGG < 1.2 ACCCCTCCCA

GAACACCAAG TGCCGAGCCT

CTGTGCTCCG GGTTCGAGG CAGCTGAGGG

CCCAAGCTY;C

CACACGAGGC

BOI TACTUCCATG AGGATGGCCA GTGGGCGAA CAUCUGGTCA CGGGTGCAG ATUACUGCAC TCCTACCGGT CACCCGGGTT GTCGCCAGT GCCGAGTC

C

<

>

0

I

0

E.

ţ,

2 1 0

CTTCTCCTTC

CTCCACTCCC

×

Z

O

M

OGATCAGCCG TCTCCCAGTG

TAACACCATT ATTYTETAA

CHACCAGCCA TECCEAGECA ATARECACTE GAUGGTERS ACRESTERING

901 GTGCCCAGGG CACCTTCAAG CCCCTGTCAG GAGAAGGCTC
CACGGGTCCC GTGGAAGTTC GGGGACAGTC CTCTTGCCAG

ACCCCTCCCT

0

ţ,

د

<u>-</u>

×

Ŀ

۲

Ç

0

<

272

AGACGGTCAC

CCTACTCGGC

H

Ξ

S

30/35

CICTCCTCTC

CCCACCCGCA

GCCOOCAGTO

CCTCACCTAC GCCCTCCGCT

CCCCAGACCA

TOCCCCCCTG GAGTCTGGTG

۲.

338

1101

CTTGCCGAGG

CAACCCCTCC

TTTCCCCCCT

TTCGCCTCCG CCGAGCCTCO

AAAGGGCCCA

GCCTCCCACC

AAGCCGAGGC

CCTCCCCACC CCACCCTCC

CCACCCTGCA

1001 CCCCCTCGGG TACTTCCGGG CACGCACAGA CCCCCGGGGT

CHECKENICE

308

ATCAAGGCCC GTGCGTGTCT GGGGGCCCCA

<

CCTCCCACCT

Ų

~

CCTATACCTT CCGAGGACAC CCTGACTTCA **GCCTCGCCCT** Œ ACCCTACCT TCCCCATGCA COCCCTCAC U ACCACCAAGC TUTTOCTICG CGGGAAGGGGAA < GCACTCCATG GGACCTGGTG GAGCCCTCGG CTCGGGACCC CCTCGACCAC CGGCTCTCCT ວນນນນນວວນນ CTCAGACCAC TCCCTCCACC TCGAATCGAG TCCCCCCCTCACCACCTCG ACCTTACCTC ACCCCCCCACC TGAAAACTGG < CCCTCTGGAC Ŋ 3 ŭ

GCATATGCAA Y T F GGACTGAAGT P D F T Œ Ç **=** > > > 3 Œ 1201 CCCCCTGCGG GGGAGCCTG ACTITITICACC CCGGCCCCCGG CCCCCACCCC 372

TCCTCCAGTC AGGACGTCAC CTCTCCATGG CACACCTACC TOCTCACTOG ACCACTGACC ۵ TCTCAATGTC ACACTTACAG z > CTAMACTCCC CATTITICAGCC ۵ w כנכנננננכענפ CCCCCCTCC > CTTAGCCACG ACTECAGTGA CGTAACTTGC CCCATAGGAG GAATCGGTGC TGAGGTCACT GCATTGAACG GGGTATCCTC ပ z < 1301 40%

GCTGGACTAC GAGGTCAAAT CTCCACTTTA CCACCTCATG U. U) Œ :: 4) ก

.....

FIGURE

CCACCTACCT
CGTCGATCGA
S Y L

AAGCGGGGAG TTCGCCCCTC

CCCCCCCCAC 'R G L

CCCCTCTCCA

TCAGAAAACC AGTCTTTTGG S E N R

¥

ACCATGAGAA GGGGGCGCGAG GGTCCCAGGA GGTTGCGGTT CCTGAAGACG TGGTAGTGTT CCCGCGCGCTC CCAGGGTCGT CGTACGGTACTTCTGC H E K G A E G P S S V R F L K T

<

C

472

1501

8C

FIGURE

CCCCCTCGTC
R E 0

COCHOOCTO

CHCCAATGAGA

CCAGACCCAA

AACATCACAG TTGTAGTGTC

AAGCCGCAGG

1601 CONCRECAT COCCCCCT CTCAGGCCCC CTACCACCC

<

Œ

<

=

O

505

O

政

Ø

0

د

0

0

S

=

0

AATCCCACACAC
TTACCCTCTC
N G R E GEOCGAGTIC TCTOCCTCAD DAAOCADAGC cricorcico W 0 K CAGCOTCAAO AOACOOAOTC u > CCTCATTOTC CCAGTAACAC > > Transcencer ACCAGGACCA >. > CAGG 1701 CTGGCCCTGA TTGCGGGGAC GGCAGTCGTG GGTGTCGTCC GACGCCCGTG CCGTCAGCAC CCACACCAGG > 5.38

GACCCTAATO ACRICTOTOAG TCCCACACTC CTCCCATTAC z ۵ CACTTATCA ACCTGGGGAA GTGAATACTT H TCCACCCCTT Ŀ AACCTCTACA TTCCAGATGT > ĸ TRITACCATICA ACATGGTACT Ξ CACCGACAGT ATCTCATCGG TTCGTCTTAT AAGCCTGTTT GTGCCTGTCA TAGAGTAGCC 0 C = TTCCCACAA ĸ **=** NACCAGNATA < 1801 572

CARGCCCCA GTTCCCCCCCT aggagagact CCCCCCCCO K 0 CTCCACACOD Œ TCAGTTTGGC GAGGTGTGCC ACTCAAACCG Ü W TTOCTCCAGG AACCACGTCC Ö O GAAGAGGTGA CTTCTCCACT > w CATT CTM 1901 GGAATTIGGA AAAGAGATCG ATGICTCCTA CGTCAA CCTTAAACGT TITCTCTAGC TACAGAGGAT GCAGTT K > > Ç, ۵ ĸ ٤. 605

ATCCCCCACT TACCCCCTCA 0 U **OCCUPATION** CCCCACCTAG TTCTOACCOA AACACTCCCT P) S د CCCCCACTCA COCCCTCACT ~ α CCACCCCCAC CCTCCCCCTC 0 Œ ts: CACCGATGTC CINCCITACAC H C 2001 GCGAAGAARS AGAGCTGTGT CACAATCAAG ACCCTGAAGG TKICGACTTCC C TCTCGACACA CCGTTAGTTC < > υ H. CCCTTCTTCC ¥ × 6.38 G

AACOCCCCC TOGACTCCTT
TTOCCCCCCC ACCTGAGGAA ī ٥ < v z **GTTCATCXA** CAAGTACCTC ĸ I ٤. AAGAGTGTCT TTCTCACAGA ы . T CCCCTCATCA COCCAGTACT I > ۵ CAACAGCATG CTTCTCCTAC I (1) z 2101 TCGAGCACCC CAATATCATC CGCCTCGAGG GCGTGGTCAC = 672

FIGURE 8D

GARCTACOTO AACCCCTCTA TTOCCCAGAT I 20 < TACCCCATCC ATXXXXTACC TECENOGRAT CONCITORAGE ACCENTAGE CONTRACTOR ACCENTAGE CONTRACTOR ACCENTAGE A Ç, CCTGCCCGAT TYCCTGCCTG TCAAGTCTCA CTACCAGCTC GTCCCCGTTCC Ü ۵ z 2201 705

AACTCTTCCC TTCAGAAGCC COACCTCCTC CCTOCAGGAG TTTCCCOATT AAARGGCTAA S R P GACTTTGGCC CTOAAACCCG ۲ C GTTTCACAGA K V S CAMAGRICATOR ACCTCGTCTG TUCAGCAGAC GTCAACAGCA Z CAACATCCTA GTTGTAGGAT z 2301 CACCGAGACC TGGCTGCTCG GTGGCTCTGG ACCGACGAGC ¤ = 7) A

GTGATGCCTG CACTACGGAC TCAACCCCCT ACTROCOCCA CCCCAAGTIC <u>~</u> CCATTCCCTT CCGATGGACT GGCTACCTGA R W T AGATTCCCAT TAGGGTA CTCCCACCAA CACCCTCCTT L G G K CACCACCTCC CTGCTCCAGG ATCCCACCTA TACKICTCICAT 2401 172

ACAGGACTAC TCTCCTCATC 0 ATGCCATTGA < GACCTGATCA CTCCACTAGT GAGCAATCAG CTCGTTAGTC 0 Z ທ ACTOGGACAT TCACCCTCTA I ۵ I GTCATTTGGG GAGAGGCCGT CTCTCCCCCA **>** _ ~ Œ, CAGTAMACCC TAACACTACA CCCTCCACTA 2501 GAGTTACGGG ATTGTGATCT GOGAGGTGAT I > CTCAATGCCC 808

CCCGTCCACC CCCCAGGTCC 0 CCCCCCTTC CCCCCCAAC ۵, CAGAAAGACC GGAATGCCCG CCTTACGGGC z CTCTTTCTCG 0 CCACTCTTCC CCTGACAACC _ 2601 CGGCTGCGC GGCCCCCAGA CTGTCCCAGC TCCCTCCAGC AGCTCATGCT GCCGGGGGTCT GACAGGGTCG AGGGAGGTCG TCCAAGTAGGA TCAAGTAGGA AGGGAGGTCG TCCAAGTAGGA TCAAGTAGGA AGGGAGGTCG TCCAAGTAGGA TCAAGTAGGA AGGGAGGTCG TCCAAGTAGGA TCAAGTAGGA TCAAGTAGA TCAAGTAGGA TCAAGTAGA TCA

TCCCCCTCCC CTCCTOOACC AGCGGCAGCC 0 GACCACCTRIC ۵ _3 GACTICTICICA CTCACACCCT ے = S TACCCCCCC CAAAATCGTG GCCCGGGAGA ATGCCGGGC < Ç v COCCCCTCT Z ĸ = CTTTTAGGAG > × CCCCCAGCCT CCCCCTCCCA 2701 TCAGCGGCT CCACAAGATG ATCCGGAACC TACCCCTTCC CCTCTTCTAC I ACTY: GCGGGA R_{J}^{2}

CAGGAAGCTC S F E CTCCTTCGAG Trescadece erocertres GACCCANACC O AAGCGTCGCC < OCT TOTAL CCAACAACT S M Þ GCCATCAAAA TXXXAAGATA COUTAGETTE ACCCETACTAT **>** = O I השממרדדכמם CACCCAACCC 2801 TUNCTACTOR GOTTTTGGCT CTGTGGGCGA
AUTUATGAGT CGAAAACUGA GACACCGGT
905 11 Y S A F G S V G E 905

CTCCTCTCAC GACCACAGTG

TCCCAACTTG ACCCTTCAAC

AAAATCATTG GOGTTTGTAG

GCCCAGTGAC CCCCTCACTC

TTTTAGTAAC

۲

۵

O

0

U

O

CCCACAGTCC

CCCCTTCTTC

GGTCGAGGTC TCACCCCCC GACAGGTCC

Ç

ບ

S

0

1105

CCCAAACATC

U

8 E FIGURE

ATGAAGTCCC TACTTCAGGO TOTOCAGCAC ACAGGTCGTC 0 > TCTTGGCCAG AGMACCOCTC Ŋ CAGAAGAAA CTCTTCTTT K 0 CCCCCCACAC CCCCCTGTG = CTCCGAATCG GAGTCACTCT CTCAGTGAGA GAGGCTTAGG _ 0 918

CATTITICCCG GTAMAACCC Accoccance 0) < CCCCAGGGAC осостесств ۵ TGACCTGCAG GAACTCCCCA CTTCACCCCT ACTIXIACGIC GGGCGTCATG CCCGCAGTAC 0 CAGGACCGGC כדניכדנונונינים 1001 AGGCCAAGCC GGAAGCCCG GGTGGGAGAG TUCGGTTCGG CCCTTCGGGC CCACCCTGTC C Ö ⊢ ပ 972

GACAGGATTT AATTTOCAGA TTAMCCTCT CCCTCGCGTC ACCAGTTCGC TCCTCAACCO O * S **E** GCCACCCCAC' > O ď CACTITICAGE 0 ت CCCTCGATTG CCCACCTAAC CCCTGTGCCC CCCACACGGG 1101 GGCAGAGTOG GGACTCACAG AGGCCCCCAG CGGTCTCACC CCTGAGTGTC TCCGGGGGTC < E. _ < 1005

TEGETETEAC ACCCACACTG CCCTCAGGAC CCCACTCCTC ۵ 0 GCCCCCCTTT CCCCCCCCAAA GAACTCCAGA CCAAGGGTGA CITTCCCACT CTTCACCTCT tr: z CCACCTCGGG U CACCCCCAG GGGGGTTCTG CCATAATAGG AGGGGAAAAT TCCCCTTTTA CCCCCAAGAC GGTATTATCC Ç --< v: 1201 1018

CTGAGGGAAC GACTCCCTTG TTCTCTCACA AAGAGAGTGT CCCCAGACCA GCCSTCTOCT ۵ ¤ **ACCITICATIVE** CCCACCCAAG v **=** ACTCCAACTA S P O W TCACCTTGAT COTCCCCCC CCACGGGGG > AGCCTCCCCA TCCCAGGGGT 0 AACATCTCCC CCAAGTCCCC CCTTCACCCCC 3301 CAGAGGAMA GTCTCCTTTT K 1072

CCCTCTCACC CCCCAACAAC 3401 CCAGUTCCAG AGTGGGGGG CTGTCCCAGG

TATTETE THEORY TTCACTTTTO TGCCTTCATA TTGAAGGTTT ACCOMMITAT AATGCCCCTC CCCCAGCTGC GGGCTCGACG TTACGGGGAG 1501 CACCAAACTC AATCATTTTT TTCCCTTGTA
GTGGTTTGAG TTAGTAAAAA AAGGGAACAT 1138 11

ATTANAAAGA O F F S AAAAACCAGA S 3 AACTCAAAAC U AACTTCCAAA

FIGURE 8F

AACTITICTOT TGGAGGGAAC CTGTTTCACT ATOGCCTCCT TTGCCCAAGT TTGAAACACA ACCTCGCTTG GACAAAGTGA TACCGGAGGA AACGGGTTCA B F V L E G T C F T M A S F A Q V > TCCTTGTCAT AGGAACAGTA CCCCGTTCCC TTITICTTIC TTCGTTTTGT TTTTCTACCG GCCCCAAGGG AAAAACAAAG AAGCAAAACA AAAAACATGC 1172

CCCCACCCCG CCTCCACCC CCAAGCTGTG TCCTATGAAGCCCTCACCCC CCACCCTCCACAC AGGATACTTC = TCCCACATCC AGGGTGTAGG CCCTTCCTCA CCCAACCAGT 1701 TGAAACAGG GCCCATCATC ATGTCTGTT CCAGAACAGT ACTTTGCC CGGGTAGTAG TACAGACAAA GGTCTTGTCA 1205 E T G A H H H V C F O N S 1205

TAAAAAAGTA Z TEGAACCCAG AAACGGACGC COGTGCTTNG AGAGGTACTT AAATTATATTA
ACCTTGGGTC TTTGCCTGCG GCCACGAACC TCCCCAAGAA TITAATATAA
E P R N G R R C L E G F L N T I JROI GGGTGTGGG TGAGGTAGTG AAAAGGGCGG TAGTTGGTGG CCCACACCCC ACTCCATCAC TTTTCCCGGC ATCAACCACC

ARNILVNSNLVCKVSDFGLSRFLEDDTSDPTYTSALGGKIPMRWTAPEAIQYRKFASAS

FIGURE 9

NVLVKSPNHVK I TDFGLARLLEGDEK EYNADGGKMP I KWMALECI HYRKFTHQS	NCMLAGDMTVCVADFGLSWKIYSGATIVRGCASKLPVKWLALGSLADNLYTVHS	NCLVGKNYTIKIADFGMSRNLYSGDYY	TRNILVENENRVKIGDFGLTKVLPQDKEYYKVKEPGESPIFWYAPESLTESLFSVASD	ARNILVNSNLVCKVSDFGMSRVLEDDPEAAYTTRGGKIPIRWTAPEAIAYRKFTSASD
FIGURE 10	FIGURE 11	FIGURE 12	FIGURE 13	FIGURE 14

1. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) According to International Patent Classification (IPC) or to both National Classification and IPC Int.C1. 5 C12N15/12; C12N15/54; C12N9/12; //C12Q1/68, C12N15/11 II. FIELDS SEARCHED Minimum Documentation Searched? Classification System Classification Symbols Int.Cl. 5 C12N; C12Q; **CO7K** Documentation Searched other than Minimum Documentation to the Extent that such Documents are included in the Fields Searched III. DOCUMENTS CONSIDERED TO BE RELEVANT Citation of Document, 11 with indication, where appropriate, of the relevant passages 12 Category * Relevant to Claim No.13 **NEURON** 1-7 vol. 6, no. 5, May 1991, pages 691 - 704 LAI, C. & LEMKE, G. 'An extended family of protein-tyrosine kinase genes differentially expressed in the verterbrate nervous system' see the whole document Special categories of cited documents: 10 later document published after the international filing date "A" document defining the general state of the art which is not or priority date and not in conflict with the application but cited to understand the principle or theory underlying the considered to be of particular relevance invention earlier document but published on or after the international "X" document of particular relevance; the claimed invention filing date cannot be considered novel or cannot be considered to "L" document which may throw doubts on priority claim(s) or involve an inventive step which is cited to establish the publication date of another "Y" document of particular relevance; the claimed invention citation or other special reason (as specified) cannot be considered to involve an inventive step when the "O" document referring to an oral disclosure, use, exhibition or document is combined with one or more other such docu-Other means ments, such combination being obvious to a person skilled document published prior to the international filing date but in the art. later than the priority date claimed "A" document member of the same patent family IV. CERTIFICATION Date of the Actual Completion of the International Search Date of Mailing of this International Search Report 0 2 -07- 1993 07 JUNE 1993 International Searching Authority Signature of Authorized Officer ANDRES S.M. EUROPEAN PATENT OFFICE

Form PCT/ISA/210 (second sheet) (Jennary 1965)

1-7, 13-15
1-7, 13-15
1-7, 13-15
13-15
13-15
1_7
1-7
1-7, 16-18
1-12

Form PCT/ISA/210 (extra sheet) (Jensery 1985)

TI

B

2.

. B

Τ.

3.

4

.

F٠